



STIC Search Report

Biotech-Chem Library

STIC Database Tracking Number: 116705

TO: Dwayne C Jones
Location: REM-4C70-4a71
Art Unit: 1614
Wednesday, March 24, 2004

Case Serial Number: 10/603677

From: Barb O'Bryen
Location: Biotech-Chem Library
Remsen E01A69
Phone: 571-272-2518

BSB
barbara.obryen@uspto.gov

Search Notes

1 of 25 102 b for
for compound, claim 3

7 view
not and
6,620,868 for wolf
6,624,177 for no off
6,624,177
13 of 25 5,948,780 for w. oce
14 of 25 3,801,633
3,850,968

2 of 25
for compound or claims
for claim treaty art 16 of 25
info

15 of 25 (compound)
16 of 25 (compound)
17 of 25 (3,850,968)
compound only

3 of 25

5 of 25
(view 5,948,780
for general treaty of NMP inhibition

6 of 25
(view 6,294,671)
general treaty of NMP inhibition

23 of 25 ?

116705

Access DB# _____

SEARCH REQUEST FORM

Scientific and Technical Information Center

U.S. Patent and Trademark Office

Requester's Full Name: Ducayne C. Jones USA Examiner #: 7099 Date: 11 MAR 04
 Address: 1614 Phone Number: (404) 557-0571 Serial Number: 10/63672
 Mail Box and Bldg Room Location: Rem 4c70 Results Format Preferred (circle): PAPER DISK E-MAIL

If more than one search is submitted, please prioritize searches in order of need.

Please provide a detailed statement of the search topic, and describe as specifically as possible the subject matter to be searched. Include the elected species or structures, keywords, synonyms, acronyms, and registry numbers, and combine with the concept or utility of the invention. Define any terms that may have a special meaning. Give examples or relevant citations, authors, etc. if known. Please attach a copy of the cover sheet, pertinent claims, and abstract.

Title of Invention: We Entangle a Sheet

Inventors (please provide full names): 11

Earliest Priority Filing Date: 11

For Sequence Searches Only Please include all pertinent information (parent, child, divisional, or issued patent numbers) along with the appropriate serial number.

Please search the compound of
 claim 3 and then crossing that
 w/ treating MS (claim 1)
 and arthritis (claim 2)

STAFF USE ONLY
 Searcher: JMTPB
 Searcher Phone #: _____
 Searcher Location: _____
 Date Searcher Pick-up: 3-24-04
 Date Entered: 3-24-04
 Searcher Prep & Review Time: 20
 Clerical Prep Time: _____
 Total Time: 9

Type of Search	Vendors and cost where applicable
NA Sequence (#)	STN <u>305</u>
AA Sequence (#)	Dialog _____
Structure (#)	Quicklit, Oram _____
Bibliographic	DB Link _____
Litigation	Lexis, Nexis _____
Fulltext	Sequence Systems _____
Patent Family	WWW, Internet _____
Other	Other _____

\$%^STN;HighlightOn=;**HighlightOff=**;
 => fil reg; d stat que 17; fil capl uspatf toxcenter; s 17
 FILE 'REGISTRY' ENTERED AT 14:32:27 ON 24 MAR 2004
 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
 PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
 COPYRIGHT (C) 2004 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

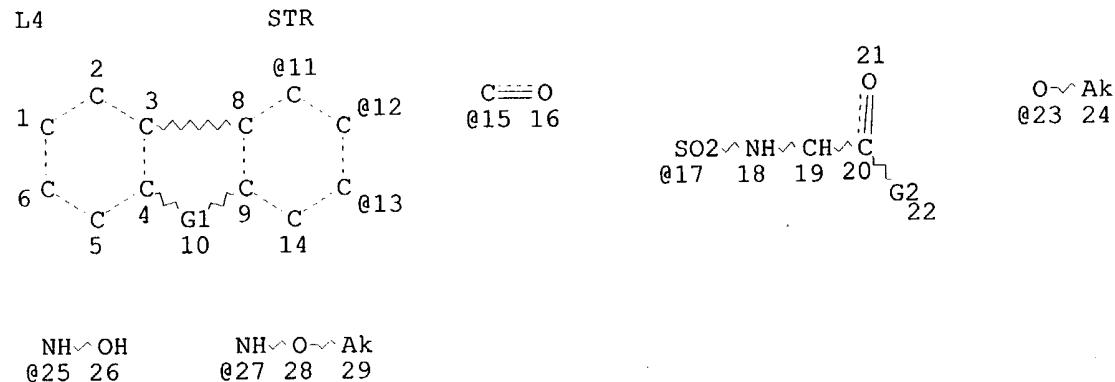
STRUCTURE FILE UPDATES: 23 MAR 2004 HIGHEST RN 666817-09-0
 DICTIONARY FILE UPDATES: 23 MAR 2004 HIGHEST RN 666817-09-0

TSCA INFORMATION NOW CURRENT THROUGH JANUARY 6, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:
<http://www.cas.org/ONLINE/DBSS/registryss.html>



VAR G1=S/CH2/15/N
 VAR G2=23/OH/25/27
 VPA 17-11/12/13 U
 NODE ATTRIBUTES:
 CONNECT IS E1 RC AT 24
 CONNECT IS E1 RC AT 29
 DEFAULT MLEVEL IS ATOM
 DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
 RING(S) ARE ISOLATED OR EMBEDDED
 NUMBER OF NODES IS 28

STEREO ATTRIBUTES: NONE
 L7 69 SEA FILE=REGISTRY SSS FUL L4

100.0% PROCESSED 2003 ITERATIONS
 SEARCH TIME: 00.00.01

69 ANSWERS

FILE 'CAPLUS' ENTERED AT 14:32:28 ON 24 MAR 2004
 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
 PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
 COPYRIGHT (C) 2004 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'USPATFULL' ENTERED AT 14:32:28 ON 24 MAR 2004
 CA INDEXING COPYRIGHT (C) 2004 AMERICAN CHEMICAL SOCIETY (ACS)

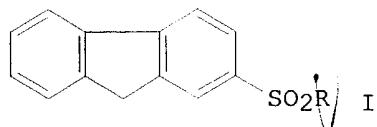
FILE 'TOXCENTER' ENTERED AT 14:32:28 ON 24 MAR 2004
 COPYRIGHT (C) 2004 ACS

L10 28 L7

=> dup rem 110
 PROCESSING COMPLETED FOR L10
 L11 25 DUP REM L10 (3 DUPLICATES REMOVED)
 ANSWERS '1-17' FROM FILE CAPLUS
 ANSWERS '18-23' FROM FILE USPATFULL
 ANSWERS '24-25' FROM FILE TOXCENTER

=> d ibib ed abs hitstr 1-23; d iall 24-25

L11 ANSWER 1 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 1
 ACCESSION NUMBER: 1991:43546 CAPLUS
 DOCUMENT NUMBER: 114:43546
 TITLE: Synthesis of biologically active fluorene-2-sulfonylamino acid and dipeptide derivatives
 AUTHOR(S): Abdel-Ghaffar, S. A.; Abbas, Y. A.
 CORPORATE SOURCE: Fac. Sci., Al-Azhar Univ., Nasr-City, Egypt
 SOURCE: Journal of the Serbian Chemical Society (1990), 55(6),
 311-17
 CODEN: JSCSEN; ISSN: 0352-5139
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 114:43546
 ED Entered STN: 09 Feb 1991
 GI

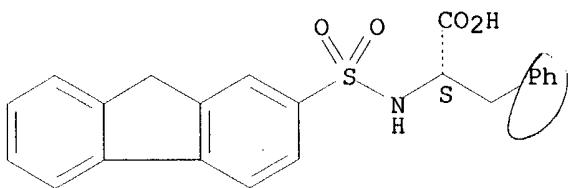


AB The prepn. of several new title derivs. I (R = X-OH, X-OMe, X-NHNH2, 2,4-C12C6H3NH, 2,4-Br2C6H3NH; X = Ala, Val, Leu, Phe, Ala-Val, Val-Leu, Leu-Ala, Ala-Phe Val-Phe) is described. Eighteen I were active against a no. of microorganisms.

IT 40356-17-0P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (prepn., esterification, and bactericidal activity of)

RN 40356-17-0 CAPLUS
 CN L-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



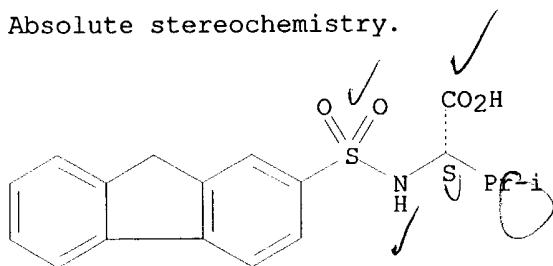
IT 32945-11-2P 56211-81-5P 131520-62-2P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (prepn., esterification, peptide coupling, and bactericidal activity
 of)

RN 32945-11-2 CAPLUS

CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

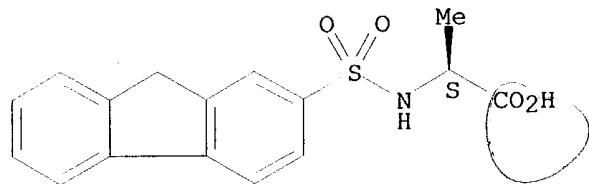
Absolute stereochemistry.



RN 56211-81-5 CAPLUS

CN L-Alanine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

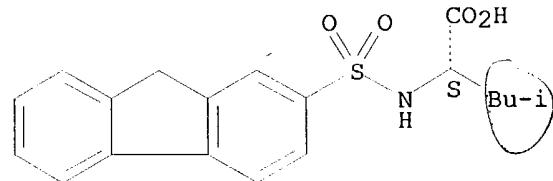
Absolute stereochemistry.



RN 131520-62-2 CAPLUS

CN L-Leucine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



IT 131520-69-9P 131520-70-2P 131520-71-3P

131520-72-4P

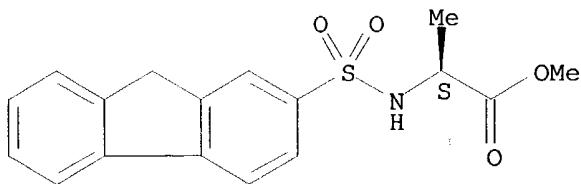
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
 (Reactant or reagent)
 (prepn., hydrazinolysis, and bactericidal activity of)

RN 131520-69-9 CAPLUS

CN L-Alanine, N-(9H-fluoren-2-ylsulfonyl)-, methyl ester (9CI) (CA INDEX

NAME)

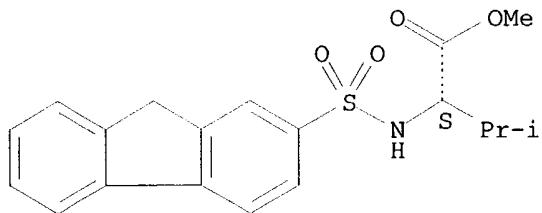
Absolute stereochemistry.



RN 131520-70-2 CAPLUS

CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)-, methyl ester (9CI) (CA INDEX NAME)

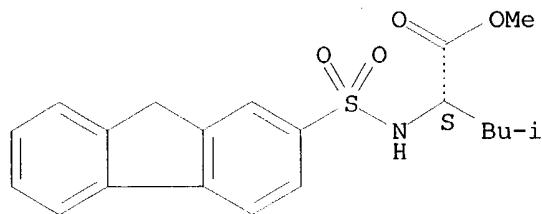
Absolute stereochemistry.



RN 131520-71-3 CAPLUS

CN L-Leucine, N-(9H-fluoren-2-ylsulfonyl)-, methyl ester (9CI) (CA INDEX NAME)

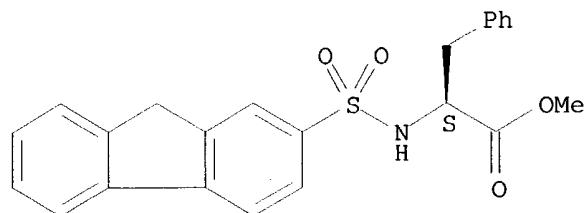
Absolute stereochemistry.



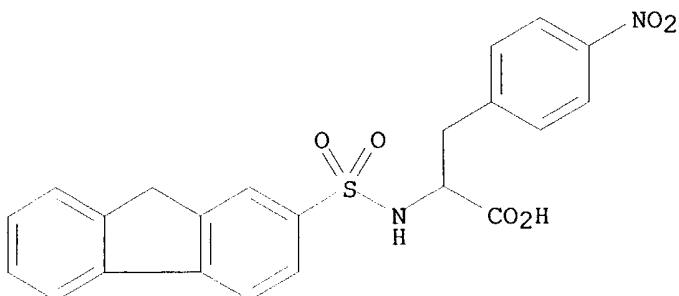
RN 131520-72-4 CAPLUS

CN L-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-, methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

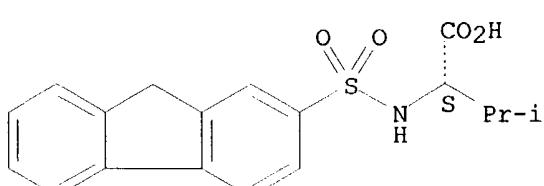


ACCESSION NUMBER: 1975:508314 CAPLUS
 DOCUMENT NUMBER: 83:108314
 TITLE: *Syntheses of amino acid derivatives and their biological activities. I. Antiinfluenza activity*
 AUTHOR(S): Kanao, Seizo; Toyoda, Takeshi; Suyama, Tadashi;
 Toyoshima, Shigeshi
 CORPORATE SOURCE: Cent. Res. Lab., Ajinomoto Co., Inc., Kawasaki, Japan
 SOURCE: Yakugaku Zasshi (1975), 95(4), 397-401
 CODEN: YKKZAJ; ISSN: 0031-6903
 DOCUMENT TYPE: Journal
 LANGUAGE: Japanese
 ED Entered STN: 12 May 1984
 GI For diagram(s), see printed CA Issue.
 AB Among 325 amino acid derivs. tested for antiviral activity, 39 of them had some activity, while the following 5 had appreciable activity:
 N-benzyl-L-valine [15363-84-5], N-furfuryl-L-phenylalanine [33014-71-0],
 N-furfuryl-4-nitro-L-phenylalanine [40356-14-7], N-2-fluorenesulfonyl-.beta.-alanine (I) [32869-90-2], and N-.beta.-naphthylaminomethyl-L-alanine [32945-07-6]. These compds. were effective when administered to mice even 72 hr after viral infection. I had both antiviral and antiinflammatory activities. The synthesis of 7 amino acid derivs. are described.
 IT 32925-03-4 32945-11-2 56211-81-5
 56211-82-6 56211-83-7
 RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)
 (antiviral activity and toxicity of)
 RN 32925-03-4 CAPLUS
 CN Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-4-nitro- (9CI) (CA INDEX NAME)



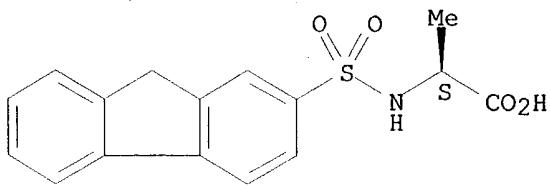
RN 32945-11-2 CAPLUS
 CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 56211-81-5 CAPLUS
 CN L-Alanine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

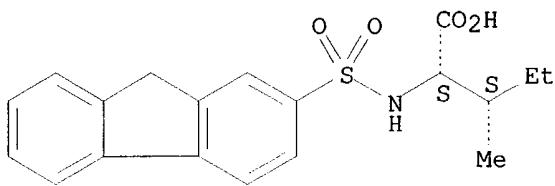
Absolute stereochemistry.



RN 56211-82-6 CAPLUS

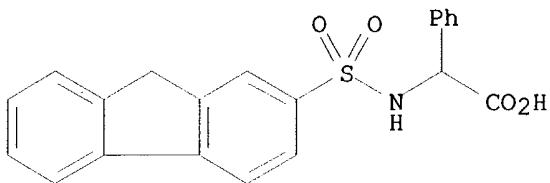
CN L-Isoleucine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 56211-83-7 CAPLUS

CN Benzeneacetic acid, .alpha.-[(9H-fluoren-2-ylsulfonyl)amino]- (9CI) (CA INDEX NAME)



L11 ANSWER 3 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 3

ACCESSION NUMBER: 1973:136657 CAPLUS

DOCUMENT NUMBER: 78:136657

TITLE: Antiviral, antiinflammatory, and antitumoral
N-substituted amino acidsINVENTOR(S): Toyoshima, Shigeshi; Kanao, Sizo; Toyoda, Takeshi;
Suyama, Tadashi; Shimizu, Akira

PATENT ASSIGNEE(S): Ajinomoto Co., Inc.

SOURCE: Ger. Offen., 23 pp.
CODEN: GWXXBX

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 4

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 2236876	A1	19730301	DE 1972-2236876	19720727
DE 2236876	B2	19800604		
DE 2236876	C3	19810212		
JP 48028612	A2	19730416	JP 1971-63252	19710819
JP 48029723	A2	19730419	JP 1971-63250	19710819
JP 52018176	B4	19770520		
JP 48029714	A2	19730419	JP 1971-63251	19710819

NL 7211361	A 19730221	NL 1972-11361	19720818
FR 2150803	A1 19730413	FR 1972-29667	19720818
PRIORITY APPLN. INFO.:		JP 1971-63250	19710819
		JP 1971-63251	19710819
		JP 1971-63252	19710819
		JP 1971-6352	19710819

ED Entered STN: 12 May 1984

AB N-Substituted amino acids (17 compds.) were prep'd. N-.beta.-Naphthyl-aminomethyl-L-leucine, Et, N-.beta.-naphthylaminomethylearbamate, and N-(2-fluorenysulfonyl)-DL-methionine were virucidal against influenza A-2/Adachi/Tokyo 57 at .apprx.20% of their LD30. N-Lauroyl-L-leucine had antiinflammatory activity comparable to hydrocortisone.

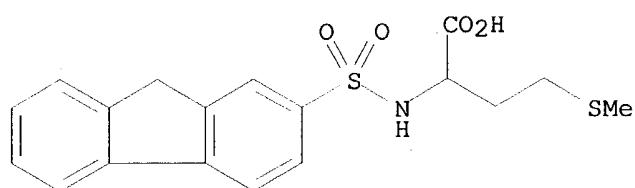
N-Ethoxycarbonylaminomethyl-DL-isoleucine, N-myristoyl-L-isoleucine, N-.beta.-naphthalenesulfonyl-DL-tryptophan, and N-propionyl-L-valine were as effective as mitomycin C at 10% of their LD30 against Ehrlich ascites and Sarcoma 180.

IT 40356-16-9P 40356-17-0P

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

RN 40356-16-9 CAPLUS

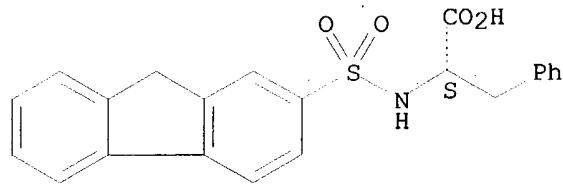
CN Methionine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)



RN 40356-17-0 CAPLUS

CN L-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L11 ANSWER 4 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2003:234336 CAPLUS

DOCUMENT NUMBER: 139:85631

TITLE: Peptides to peptidomimetics: Towards the design and synthesis of bioavailable inhibitors of oligosaccharyl transferase

AUTHOR(S): Weerapana, Eranthie; Imperiali, Barbara

CORPORATE SOURCE: Department of Chemistry, Massachusetts Institute of Technology, Cambridge, MA, 02139, USA

SOURCE: Organic & Biomolecular Chemistry (2003), 1(1), 93-99

PUBLISHER: Royal Society of Chemistry

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 139:85631

ED Entered STN: 26 Mar 2003

AB Oligosaccharyl transferase (OT) is the enzyme responsible for asparagine-linked glycosylation in the lumen of the endoplasmic reticulum, which is a subcellular compartment within eukaryotic cells. Inhibition of this enzyme within a cellular environment would provide a valuable investigative tool for glycobiol. Due to the limitations of peptides, none of the existing peptide-based inhibitors of OT demonstrate activity in cell-based enzyme assays. We report the design, synthesis and preliminary biol. characterization of a family of peptidomimetics that inhibit OT with K_i values in the nanomolar range. The hexapeptide Bz-Dab-Ala-Thr-Val-Thr-Nph-NH₂ (Dab = 2,4-diaminobenzoic acid, Nph = p-nitrophenylalanine, K_i = 69 nM) was used as the prototype for the design of bioavailable inhibitors. Several aminobenzoic acid spacer groups were evaluated as potential isosteres of the Val-Thr dipeptide unit and the peptidomimetic incorporating 3-aminobenzoic acid proved to inhibit OT with similar potency to the parent compd. (K_i = 84 nM). Further modifications explored the effects of size, hydrophobicity and conformational rigidity on enzyme affinity. This study yielded a family of potent non-peptidic inhibitors that are viable candidates for the in vivo inhibition of OT.

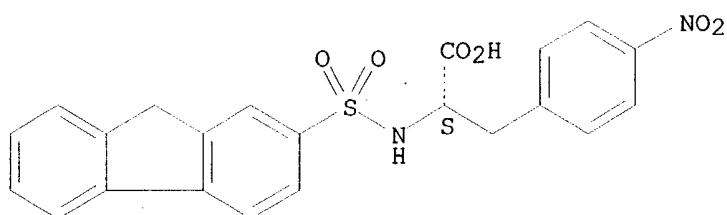
IT 52525-95-8

RL: RCT (Reactant); RACT (Reactant or reagent)
(prepn. of peptidomimetics contg. aminobenzoic acid isostere as
inhibitors of oligosaccharyl transferase)

RN 52525-95-8 CAPLUS

CN L-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-4-nitro- (9CI) (CA INDEX
NAME)

Absolute stereochemistry.



REFERENCE COUNT:

21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 5 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1998:402296 CAPLUS

DOCUMENT NUMBER: 129:76499

TITLE: Method for treating and preventing heart failure and ventricular dilation

INVENTOR(S): Peterson, Joseph T., Jr.

PATENT ASSIGNEE(S): Warner-Lambert Co., USA

SOURCE: PCT Int. Appl., 178 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9825597	A2	19980618	WO 1997-US21934	19971202
WO 9825597	A3	20001012		
W:	AL, AU, BA, BB, BG, BR, CA, CN, CZ, EE, GE, HU, ID, IL, IS, JP, KR, LC, LK, LR, LT, LV, MG, MK, MN, MX, NO, NZ, PL, RO, SG, SI, SK, SL, TR, TT, UA, UZ, VN, YU, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR,			

GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA,
 GN, ML, MR, NE, SN, TD, TG
 AU 9855906 A1 19980703 AU 1998-55906 19971202
 AU 741768 B2 20011206
 BR 9714385 A 20000516 BR 1997-14385 19971202
 EP 1028716 A1 20000823 EP 1997-952246 19971202
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, SI, LT, LV, FI, RO
 NZ 334897 A 20010223 NZ 1997-334897 19971202
 JP 2001526631 T2 20011218 JP 1998-526758 19971202
 ZA 9711004 A 19981005 ZA 1997-11004 19971208
 US 5948780 A 19990907 US 1997-987167 19971208
 NO 9902769 A 19990809 NO 1999-2769 19990608
 PRIORITY APPLN. INFO.: US 1996-32631P P 19961209
 WO 1997-US21934 W 19971202

*Different
Up*

OTHER SOURCE(S): MARPAT 129:76499

ED Entered STN: 01 Jul 1998

AB Matrix metalloproteinase inhibitors are useful for preventing and treating heart failure, and ventricular dilation in mammals. Thus, 2-(4'-bromobiphenyl-4-sulfonylamino)-3-methylbutyric acid was effective in protecting pigs in the pacing-induced heart failure model.

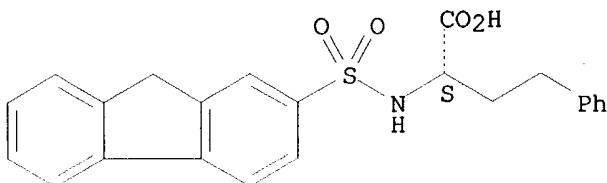
IT 204769-92-6

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (use of matrix metalloproteinase inhibitors in treating heart failure
 and ventricular dilation)

RN 204769-92-6 CAPLUS

CN Benzenebutanoic acid, .alpha.-[(9H-fluoren-2-ylsulfonyl)amino]-,
 (.alpha.S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L11 ANSWER 6 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1998:175917 CAPLUS

DOCUMENT NUMBER: 128:230699

TITLE: Preparation of dibenzofuransulfonyl and related amino acids for inhibition of matrix metalloproteinases

INVENTOR(S): Picard, Joseph Armand; Sliskovic, Drago Robert

PATENT ASSIGNEE(S): Warner-Lambert Company, USA; Picard, Joseph Armand;
 Sliskovic, Drago Robert

SOURCE: PCT Int. Appl. 83 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9809957	A1	19980312	WO 1997-US15444	19970902
W: AL, AU, BA, BB, BG, BR, CA, CN, CZ, EE, GE, GH, HU, IL, IS, JP, KR, LC, LK, LR, LT, LV, MG, MK, MN, MX, NO, NZ, PL, RO, SG, SI, SK, SL, TR, TT, UA, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				

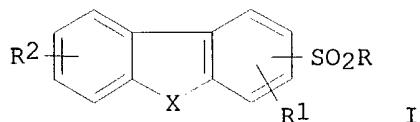
RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG

AU 9741770	A1	19980326	AU 1997-41770	19970902
AU 736347	B2	20010726		
EP 929542	A1	19990721	EP 1997-939751	19970902
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
BR 9712794	A	19991214	BR 1997-12794	19970902
NZ 333064	A	20001124	NZ 1997-333064	19970902
JP 2002514180	T2	20020514	JP 1998-512818	19970902
ZA 9707920	A	19980302	ZA 1997-7920	19970903
HR 970474	B1	20021031	HR 1997-970474	19970904
US 6294674	B1	20010925	US 1999-254403	19990302
PRIORITY APPLN. INFO.:				
			US 1996-25063P	P 19960904
			US 1997-55714P	P 19970807
			WO 1997-US15444	W 19970902

OTHER SOURCE(S): MARPAT 128:230699

ED Entered STN: 25 Mar 1998

GI



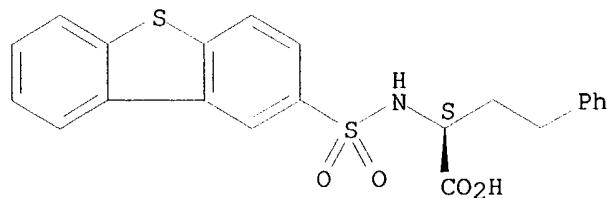
AB Heterocyclyl sulfonyl amino acids I (R = unnatural amino acid; X = O, S, SO, SO₂, CO, NH, alkyl- or alkylphenylimino; R₁, R₂ = H, alkyl, Ph, NO₂, halo, alkoxy, CN, etc.) or their pharmaceutically acceptable salts, esters, amides, and prodrugs were prepd. as matrix metalloproteinases inhibitors. Thus, 6-[2-(4-chlorophenoxy)-2-methylpropionylamino]-2-(dibenzofuran-2-ylsulfonylamino)hexanoic acid, prepd. by acylation of 6-amino-2-(dibenzofuran-2-ylsulfonylamino)hexanoic acid Me ester hydrobromide, showed IC₅₀ >100 .mu.M against MMP-1 and MMP-7.

IT 204769-50-6P 204769-91-5P 204769-92-6P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of dibenzofuransulfonyl and related amino acids for inhibition of matrix metalloproteinases)

RN 204769-50-6 CAPLUS

CN Benzenebutanoic acid, .alpha.-[(2-dibenzothienylsulfonyl)amino]-, (S)- (9CI) (CA INDEX NAME)

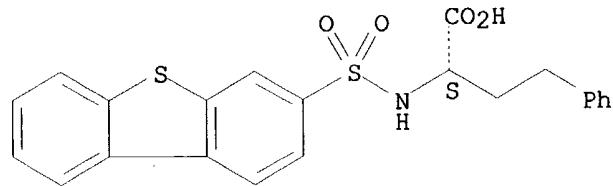
Absolute stereochemistry.



RN 204769-91-5 CAPLUS

CN Benzenebutanoic acid, .alpha.-[(3-dibenzothienylsulfonyl)amino]-, (S)- (9CI) (CA INDEX NAME)

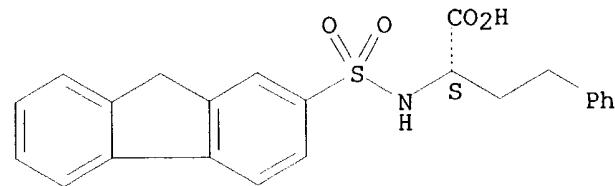
Absolute stereochemistry.



RN 204769-92-6 CAPLUS

CN Benzenebutanoic acid, .alpha.-[(9H-fluoren-2-ylsulfonyl)amino]-, (.alpha.S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 7 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1998:175896 CAPLUS

DOCUMENT NUMBER: 128:217278

TITLE: Preparation of dibenzofuransulfonamides as matrix metalloproteinase (MMP) inhibitors and their therapeutic uses

INVENTOR(S): O'Brien, Patrick Michael; Picard, Joseph Armand; Sliskovic, Drago Robert; White, Andrew David

PATENT ASSIGNEE(S): Warner-Lambert Company, USA; O'Brien, Patrick Michael; Picard, Joseph Armand; Sliskovic, Drago Robert; White, Andrew David

SOURCE: PCT Int. Appl., 58 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9809934	A1	19980312	WO 1997-US14859	19970822
W: AL, AU, BA, BB, BG, BR, CA, CN, CZ, EE, GE, GH, HU, IL, IS, JP, KR, LC, LK, LR, LT, LV, MG, MK, MN, MX, NC, NZ, PL, RO, SG, SI, SK, SL, TR, TT, UA, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
CA 2256716	AA	19980312	CA 1997-2256716	19970822
AU 9741595	A1	19980326	AU 1997-41595	19970822
AU 735013	B2	20010628		
EP 931045	A1	19990728	EP 1997-939527	19970822
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI				

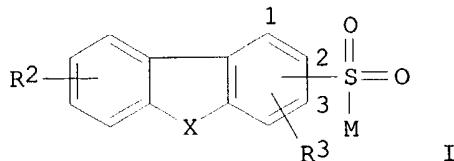
applicants' priority date

BR 9711988	A	19990824	BR 1997-11988	19970822
NZ 333063	A	20001222	NZ 1997-333063	19970822
JP 2000517341	T2	20001226	JP 1998-512709	19970822
ZA 9707916	A	19980303	ZA 1997-7916	19970903
US 6624177	B1	20030923	US 1999-254384	19990302
US 2003032665	A1	20030213	US 2002-162518	20020604
US 6620835	B2	20030916	US 2003-603677	20030625
US 2004029945	A1	20040212	US 1996-25062P	P 19960904
PRIORITY APPLN. INFO.:				
			US 1997-55713P	P 19970807
			WO 1997-US14859	W 19970822
			US 1999-254384	A3 19990302
			US 2002-162518	A3 20020604

OTHER SOURCE(S): MARPAT 128:217278

ED Entered STN: 25 Mar 1998

GI



AB A method of inhibiting MMP is claimed comprising administering to a patient a therapeutically effective amt. of dibenzofuransulfonamides [I; M = natural L-.alpha.-amino acid deriv. NHCHRCOR1; X = O, S, S(O)n, CH2, CO, NR4; R = side chain of a natural amino acid; R1 = OH, C1-5 alkoxy, NHOR5; R2, R3 = H, C1-5 alkyl, nitrophenyl, halo, cyano, etc.; R4 = H, C1-6 alkyl(phenyl); R5 = H, C1-5 alkyl; n = 0-2] including their pharmaceutically acceptable salts, esters, amides and prodrugs. A method of treating diseases in which matrix metalloproteinases are involved, e.g., multiple sclerosis, atherosclerotic plaque rupture, restenosis, aortic aneurism, heart failure, periodontal disease, corneal ulceration, burns, decubital ulcers, chronic ulcers or wounds, cancer metastasis, tumor angiogenesis, arthritis, or other autoimmune or inflammatory diseases dependent upon tissue invasion by leukocytes, and 26 specific sulfonamides are also claimed. Thus, dibenzofuran-2-sulfonyl chloride was amidated with L-leucine tert-Bu ester and the resulting tert-Bu L-2-(dibenzofuran-2-sulfonylamino)-4-methylpentanoate (64% yield) was hydrolyzed with CF3CO2H/anisole in CH2Cl2 to give 33% L-2-(dibenzofuran-2-sulfonylamino)-4-methylpentanoic acid which inhibited gelatinase A and stromelysin-1 with IC50 of 0.32 and 1.18, resp. (units not given).

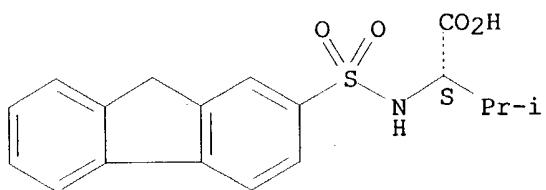
IT 32945-11-2P 204440-69-7P 204440-70-0P
204440-71-1P 204440-91-5P 204440-92-6P
204440-93-7P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of dibenzofuransulfonamides as matrix metalloproteinase inhibitors and their therapeutic uses)

RN 32945-11-2 CAPLUS

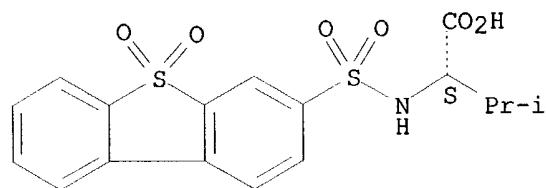
CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



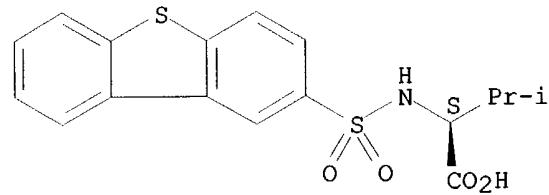
RN 204440-69-7 CAPLUS
 CN L-Valine, N-[5,5-dioxido-3-dibenzothienyl)sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



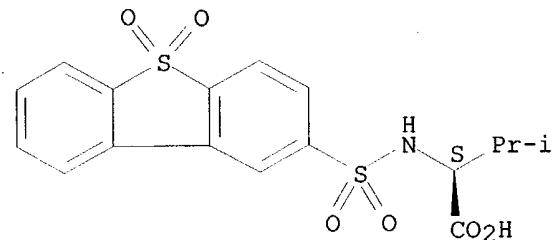
RN 204440-70-0 CAPLUS
 CN L-Valine, N-(2-dibenzothienylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



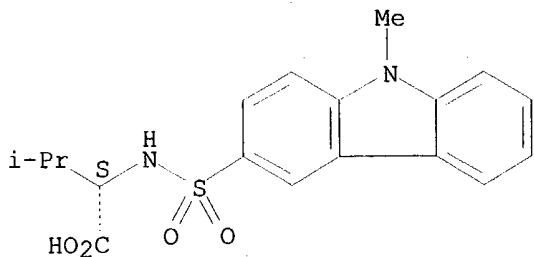
RN 204440-71-1 CAPLUS
 CN L-Valine, N-[5,5-dioxido-2-dibenzothienyl)sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



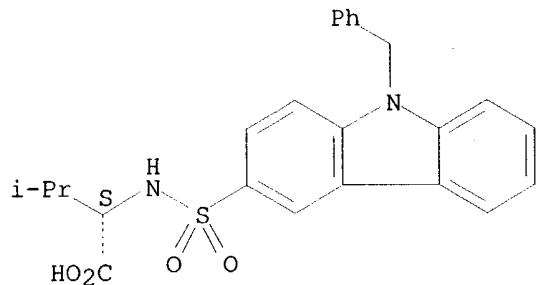
RN 204440-91-5 CAPLUS
 CN L-Valine, N-[9-methyl-9H-carbazol-3-yl)sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



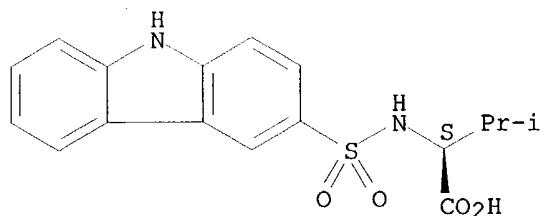
RN 204440-92-6 CAPLUS
 CN L-Valine, N-[9-(phenylmethyl)-9H-carbazol-3-yl]sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 204440-93-7 CAPLUS
 CN L-Valine, N-(9H-carbazol-3-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 8 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 1998:784844 CAPLUS
 DOCUMENT NUMBER: 130:110586
 TITLE: The synthesis and biological properties of N- and O-substituted amino acids
 AUTHOR(S): Straukas, Juozapas; Dirvianskyte, Nijole; Palaima, Algirdas
 CORPORATE SOURCE: Institute of Biochemistry, Vilnius, 2600, Lithuania
 SOURCE: Chemija (1998), (2), 160-164
 CODEN: CHMJE; ISSN: 0235-7216
 PUBLISHER: Academia
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 ED Entered STN: 15 Dec 1998
 AB The synthesis and biol. properties of nonproteinogenic amino acids derivs.

based on DL-serine, threo-DL-phenylserine, threo-DL-m-nitrophenylserine, 2-methoxy-5-nitro-DL-phenylalanine, 4-methoxy-3-nitro-DL-phenylalanine and .vepsiln.-aminocaproic acid are reported. Some derivs. of threo-DL-phenylserine exhibited antiviral activity. Thus, O-phenylacetyl-threo-DL-phenylserine Et ester hydrochloride completely inhibited the reprodn. of vesicular stomatitis virus, influenza virus A2 and type 23 adenovirus. N-(p-Brombenzenesulfonyl)-threo-DL-phenylserine Et ester and N-(2-fluorenylidene)-threo-DL-phenylserine Et ester completely stop the reprodn. of enterovirus Coxsackie A13.

IT 178633-88-0P 178633-89-1P 219642-36-1P

219642-37-2P

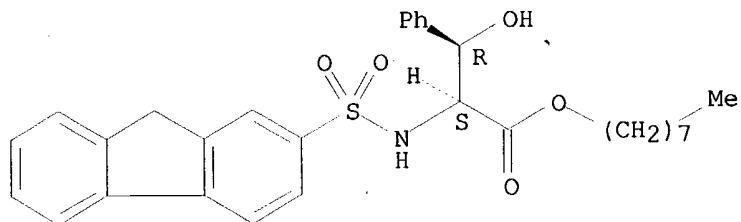
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

(synthesis and antiviral activities of N- and O-substituted amino acids)

RN 178633-88-0 CAPPLUS

CN D-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-.beta.-hydroxy-, octyl ester, (.beta.S)-rel- (9CI) (CA INDEX NAME)

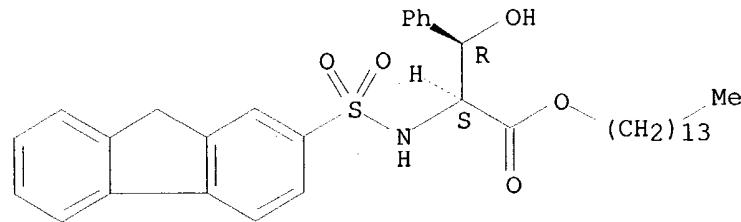
Relative stereochemistry.



RN 178633-89-1 CAPPLUS

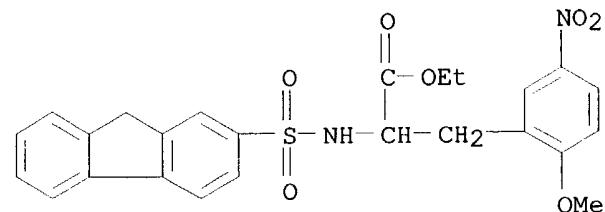
CN D-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-.beta.-hydroxy-, tetradecyl ester, (.beta.S)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

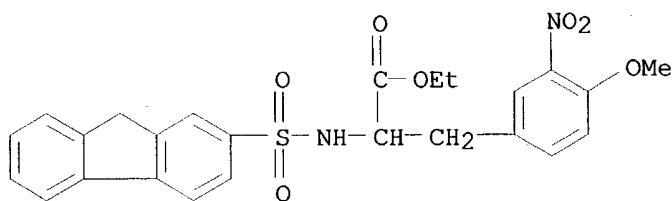


RN 219642-36-1 CAPPLUS

CN Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-2-methoxy-5-nitro-, ethyl ester (9CI) (CA INDEX NAME)



RN 219642-37-2 CAPLUS
 CN Tyrosine, N-(9H-fluoren-2-ylsulfonyl)-O-methyl-3-nitro-, ethyl ester (9CI)
 (CA INDEX NAME)



REFERENCE COUNT: 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 9 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 1996:340464 CAPLUS
 DOCUMENT NUMBER: 125:76320
 TITLE: Synthesis and antiviral activity of N- and O-substituted amino acids
 AUTHOR(S): Straukas, Yu.; Dirvyanskite, N.; Yankauskas, R.;
 Yavorovskaya, V. E.; Evstropov, A. N.; Kiseleva, V. N.
 CORPORATE SOURCE: Inst. of Biochemistry, Vilnius, Lithuania
 SOURCE: Khimiko-Farmatsevticheskii Zhurnal (1996), 30(4), 18-21
 CODEN: KHFZAN; ISSN: 0023-1134

PUBLISHER: Izdatel'stvo Folium
 DOCUMENT TYPE: Journal
 LANGUAGE: Russian

ED Entered STN: 12 Jun 1996

AB Twenty-six amino acid derivs. were prep'd. from threo-DL-phenylserine (I), DL-serine, threo-DL- α -nitrophenylserine, 2-methoxy-5-nitro-DL-phenylalanine, 4-methoxy-3-nitro-DL-phenylalanine, and .epsilon.-aminocaproic acid. Satd. carboxylic or arylsulfonic acid residues or fluorenylidene group were introduced into the mol's of amino acids and their alkyl esters. Derivs. of I had the highest antiviral activity.

IT 178633-88-0P 178633-89-1P 178633-90-4P

178633-91-5P

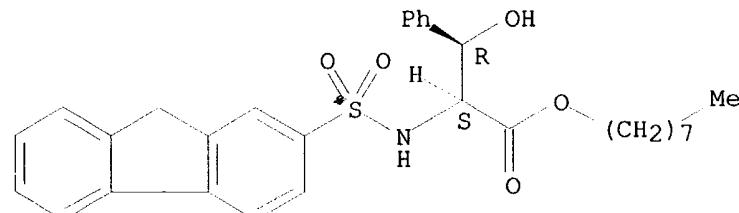
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

(prepn. and antiviral activity of N- and O-substituted amino acids)

RN 178633-88-0 CAPLUS

CN D-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-.beta.-hydroxy-, octyl ester, (.beta.S)-rel- (9CI) (CA INDEX NAME)

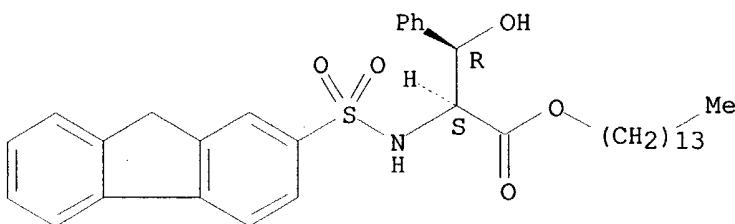
Relative stereochemistry.



RN 178633-89-1 CAPLUS
 CN D-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-.beta.-hydroxy-, tetradecyl

ester, (.beta.S)-rel- (9CI) (CA INDEX NAME)

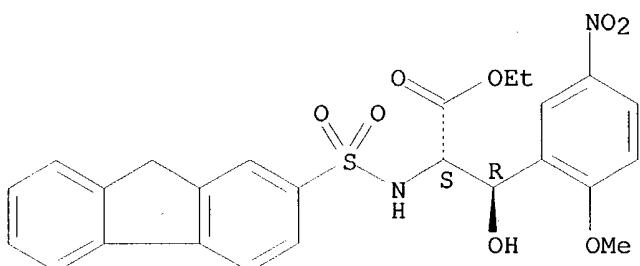
Relative stereochemistry.



RN 178633-90-4 CAPLUS

CN D-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-.beta.-hydroxy-2-methoxy-5-nitro-, ethyl ester, (.beta.S)-rel- (9CI) (CA INDEX NAME)

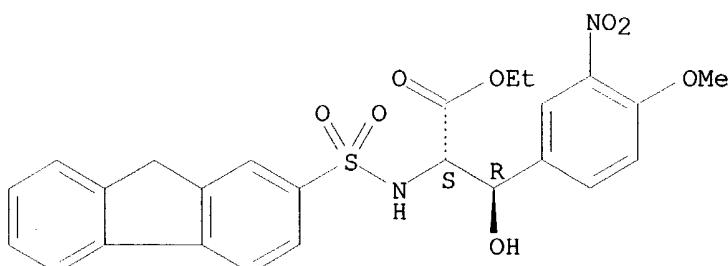
Relative stereochemistry.



RN 178633-91-5 CAPLUS

CN D-Tyrosine, N-(9H-fluoren-2-ylsulfonyl)-.beta.-hydroxy-O-methyl-3-nitro-, ethyl ester, (.beta.S)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.



L11 ANSWER 10 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1995:902595 CAPLUS

DOCUMENT NUMBER: 123:306554

TITLE:

~~N-2-fluorenonesulfonyl derivatives of
threo-DL-phenylserine showing antiviral activity with
respect to herpes simplex virus type I~~

INVENTOR(S): Straukas, Yu. Yu.; Bulko, R. E.; Yavorovskaya, V. E.;
Evstropov, A. N.; Galegov, G. A.; Pravdina, N. F.

PATENT ASSIGNEE(S): Institut Biokhimii AN LitSSR, Liechtenstein;
Novosibirskij Gosudarstvennyj Meditsinskij Institut;
Institut Virusologii im. D.I. Ivanovskogo

SOURCE: U.S.S.R. From: Izobreteniya 1993, (47-8), 173.

CODEN: URXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

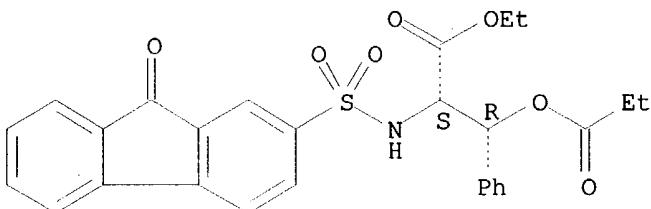
Russian

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
SU 1405269	A1	19931230	SU 1986-4128762	19860709
PRIORITY APPLN. INFO.:			SU 1986-4128762	19860709
ED	Entered STN:	08 Nov 1995		
AB	Title only translated.			
IT	169944-28-9 169944-29-0			
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)				
	(virucidal activity of fluorenonesulfonyl derivs. of threo-phenylserine against herpes simplex virus)			
RN	169944-28-9 CAPLUS			
CN	D-Phenylalanine, N-[(9-oxo-9H-fluoren-2-yl)sulfonyl]-.beta.-(1-oxopropoxy)-, ethyl ester, (.beta.S)-rel- (9CI) (CA INDEX NAME)			

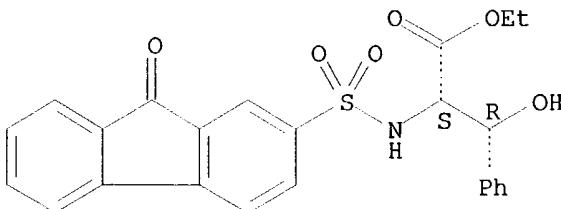
Relative stereochemistry.



RN 169944-29-0 CAPLUS

CN D-Phenylalanine, .beta.-hydroxy-N-[(9-oxo-9H-fluoren-2-yl)sulfonyl]-, ethyl ester, (.beta.S)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.



L11 ANSWER 11 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1995:902594 CAPLUS

DOCUMENT NUMBER: 123:306553

TITLE: N-2-Fluorenesulfonyl-O-propionyl-threo-DL-phenylserine ethyl ester showing antiviral activity with respect to Echo 11 virus

INVENTOR(S): Straukas, Yu. Yu.; Bulko, R. E.; Yavorovskaya, V. E.; Evstropov, A. I.

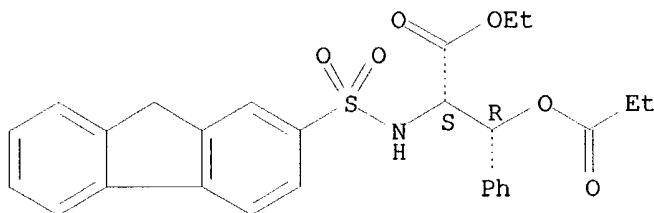
PATENT ASSIGNEE(S): Institut Biokhimii AN LitSSR, Russia; Novosibirskij Gosudarstvennyj Meditsinskij Institut

SOURCE: U.S.S.R. From: Izobreteniya 1993, (47-8), 173.
CODEN: URXXAF

DOCUMENT TYPE: Patent
 LANGUAGE: Russian
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
SU 1363764	A1	19931230	SU 1986-4015748	19860131
PRIORITY APPLN. INFO.:			SU 1986-4015748	19860131
ED	Entered STN:	08 Nov 1995		
AB	Title only translated.			
IT	169944-27-8			
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (virucidal activity of phenylserine Et ester deriv. against echo 11 virus)				
RN	169944-27-8 CAPLUS			
CN	D-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-.beta.- (1-oxopropoxy)-, ethyl ester, (.beta.S)-rel- (9CI) (CA INDEX NAME)			

Relative stereochemistry.

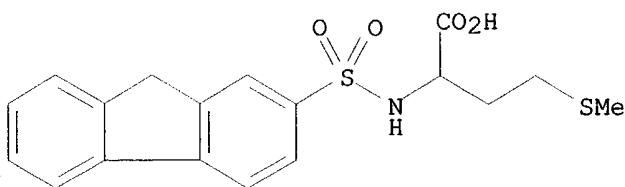


L11 ANSWER 12 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 1980:568614 CAPLUS
 DOCUMENT NUMBER: 93:168614
 TITLE: N-(2-fluorenylsulfonyl) amino acids
 PATENT ASSIGNEE(S): Ajinomoto Co., Inc., Japan
 SOURCE: Jpn. Tokkyo Koho, 3 pp.
 CODEN: JAXXAD

DOCUMENT TYPE:
 LANGUAGE:
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
JP 55004741	B4	19800131	JP 1978-105936	19780830	
JP 54048745	A2	19790417			
PRIORITY APPLN. INFO.:			JP 1978-105936	19780830	
ED	Entered STN:	12 May 1984			
AB	N-(2-Fluorenylsulfonyl) amino acid, <u>useful as viricides</u> , were prep'd. from 2-fluorenensulfonyl chloride and the appropriate amino acids. Thus, 2.7 g L-alanine in 10% aq. NaOH soln. was treated with 8.8 g 2-fluorenensulfonyl chloride at 50.degree. for 30 min to give 6.8% N-(2-fluorenylsulfonyl)-L-alanine. The N-(2-fluorenylsulfonyl) derivs. of L-isoleucine, DL-tryptophan, DL-methionine, and DL-phenylglycine were also prep'd.				
IT	40356-16-9P 56211-81-5P 56211-82-6P 56211-83-7P 61447-77-6P				
RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. of)					
RN	40356-16-9 CAPLUS				

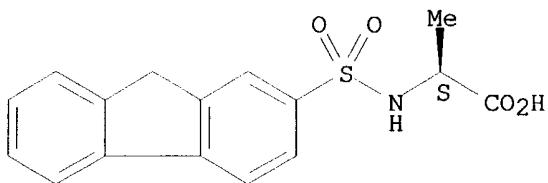
CN Methionine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)



RN 56211-81-5 CAPLUS

CN L-Alanine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

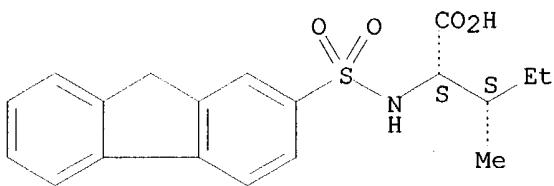
Absolute stereochemistry.



RN 56211-82-6 CAPLUS

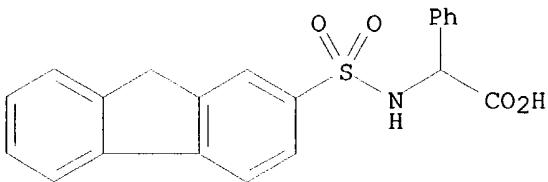
CN L-Isoleucine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



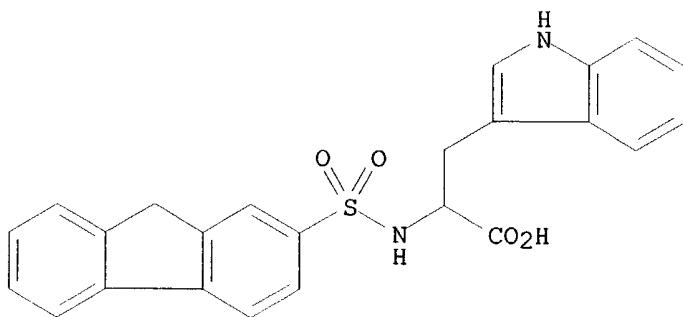
RN 56211-83-7 CAPLUS

CN Benzeneacetic acid, .alpha.-[(9H-fluoren-2-ylsulfonyl)amino]- (9CI) (CA INDEX NAME)



RN 61447-77-6 CAPLUS

CN Tryptophan, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)



L11 ANSWER 13 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1979:152610 CAPLUS

DOCUMENT NUMBER: 90:152610

TITLE: N2-Arylsulfonyl-L-argininamides

INVENTOR(S): Okamoto, Shosuke; Kikumoto, Ryoji; Tamao, Yoshikuni;
Okubo, Kazuo; Tezuka, Toru; Tonomura, Shinji;
Hijikata, Akiko

PATENT ASSIGNEE(S): Mitsubishi Chemical Industries Co., Ltd., Japan

SOURCE: Ger. Offen., 147 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 15

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 2801478	A1	19780720	DE 1978-2801478	19780113
DE 2801478	C2	19910131		
US 4066773	A	19780103	US 1977-760745	19770119
US 4073913	A	19780214	US 1977-760668	19770119
US 4093712	A	19780606	US 1977-760672	19770119
US 4097472	A	19780627	US 1977-760676	19770119
US 4101653	A	19780718	US 1977-760929	19770119
US 4097591	A	19780627	US 1977-776195	19770310
JP 54003037	A2	19790111	JP 1977-66508	19770606
JP 60010028	B4	19850314		
US 4125604	A	19781114	US 1977-804334	19770607
US 4131673	A	19781226	US 1977-804368	19770607
US 4140681	A	19790220	US 1977-804331	19770607
IL 53685	A1	19851231	IL 1977-53685	19771223
AU 7832289	A1	19790719	AU 1978-32289	19780109
AU 522320	B2	19820527		
ZA 7800123	A	19790829	ZA 1978-123	19780109
FI 7800073	A	19780720	FI 1978-73	19780110
FI 72316	B	19870130		
FI 72316	C	19870511		
ES 466706	A2	19781016	ES 1978-466706	19780110
NL 7800448	A	19780721	NL 1978-448	19780113
NL 187746	B	19910801		
NL 187746	C	19920102		
SE 7800512	A	19780720	SE 1978-512	19780117
SE 452624	B	19871207		
SE 452624	C	19880317		
HU 22709	O	19820628	HU 1978-MI626	19780117
HU 180265	B	19830228		
DK 7800263	A	19780720	DK 1978-263	19780118
DK 150521	B	19870316		

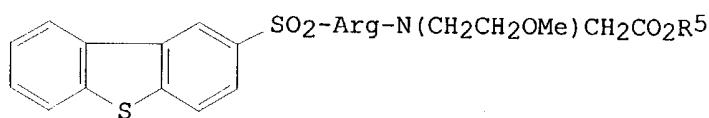
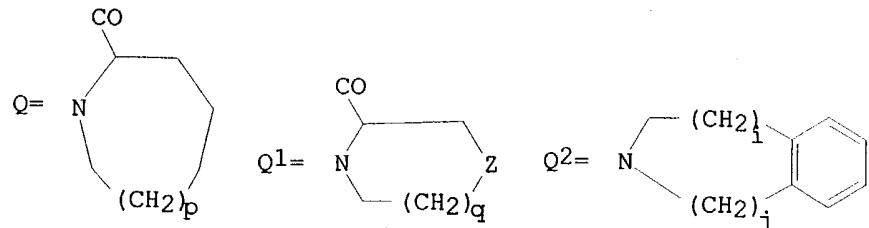
DK 150521	C	19871019		
NO 7800191	A	19780720	NO 1978-191	19780118
NO 158681	B	19880711		
NO 158681	C	19881019		
FR 2378004	A2	19780818	FR 1978-1368	19780118
FR 2378004	B2	19850913		
GB 1596971	A	19810903	GB 1978-2063	19780118
PL 123267	B1	19821030	PL 1978-204063	19780118
CH 633773	A	19821231	CH 1978-519	19780118
CH 648293	A	19850315	CH 1978-4530	19780118
SU 1181539	A3	19850923	SU 1978-2566652	19780118
BE 863092	A4	19780719	BE 1978-184463	19780119
ES 466705	A2	19790816	ES 1978-466705	19780119
DD 137352	C	19790829	DD 1978-203302	19780119
AT 7800399	A	19820515	AT 1978-399	19780119
AT 369356	B	19821227		
CS 236757	B2	19850515	CS 1978-381	19780119
JP 62014548	B4	19870402	JP 1978-4529	19780119
JP 54100342	A2	19790808		
US 4173630	A	19791106	US 1978-902855	19780504
SU 938739	A3	19820623	SU 1979-2776611	19790618
AT 8003284	A	19820515	AT 1980-3284	19800623
AT 369357	B	19821227		
AT 8003285	A	19820515	AT 1980-3285	19800623
AT 369358	B	19821227		
CS 236772	B2	19850515	CS 1981-2011	19810319
CS 236773	B2	19850515	CS 1981-2012	19810319
FI 8402539	A	19840621	FI 1984-2539	19840621
FI 74455	B	19871030		
FI 74455	C	19880208		

PRIORITY APPLN. INFO.:

US 1977-760668	19770119
US 1977-760672	19770119
US 1977-760676	19770119
US 1977-760745	19770119
US 1977-760929	19770119
US 1977-776195	19770310
JP 1977-66508	19770606
US 1977-804331	19770607
US 1977-804368	19770607
JP 1974-128774	19741108
JP 1974-128775	19741108
JP 1974-136695	19741129
JP 1974-136697	19741129
JP 1975-23268	19750225
JP 1975-23635	19750226
JP 1975-26768	19750305
JP 1975-29357	19750311
JP 1975-29358	19750311
US 1975-62	239 19751014
US 1975-622390	19751014
US 1975-638985	19751209
US 1976-646522	19760105
US 1976-649219	19760114
US 1976-653217	19760128
US 1976-656014	19760206
US 1976-656870	19760210
US 1976-669743	19760324
US 1976-671436	19760329
US 1976-671568	19760329
US 1976-703704	19760708
US 1976-707536	19760722
US 1976-713486	19760811
US 1976-723474	19760914

US 1976-728051	19760930
US 1977-760677	19770119
FI 1978-73	19780110
CH 1978-519	19780118
AT 1978-399	19780119
CS 1978-381	19780119

ED Entered STN: 12 May 1984
 GI



I

AB RSO₂-Arg-X-OR₁ [R = substituted Ph, substituted naphthyl, heterocyclic group; X = NR₂(CH₂)_nC=O (R₂ = aliph., aralkyl, carbocyclic, or heterocyclic group; n = 1-3), NR₃CHR₄(CH₂)_mC=O (R₃ = H or R₂; R₄ = C₁-10 alkyl, substituted C₁-10 alkyl, C₁-12 aralkyl, substituted benzyl; m = 0-2), substituted piperidinocarboxylic acid residue, Q (p = 1-4), Q₁ (Z = O, S, SO; q = 0, 1), Q₂ (i and j = 0-2 where i + j = 1 or 2); R₁ = H, C₁-10 alkyl, C₆-10 aryl, C₇-12 aralkyl] and their salts (.apprx.135 compds.) were prepd. as thrombin inhibitors. Thus, arginine was acylated with 2-dibenzothienylsulfonyl chloride to give the N2-sulfonyl deriv., which was converted to its acid chloride and amidated with MeOCH₂CH₂-Gly-OEt to give dipeptide I (R₅ = Et) (II). II was saponified to give I (R₅ = H) (III). III at 0.45 .mu.M doubled blood coagulation time.

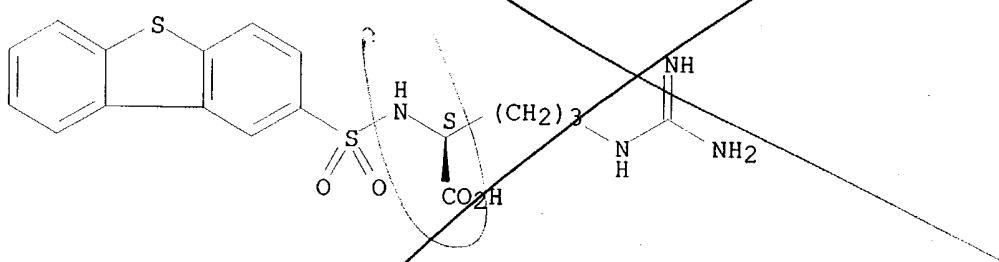
IT 69129-22-2P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (prepn. and acid chlorination of)

RN 69129-22-2 CAPLUS

CN L-Arginine, N2-(2-dibenzothienylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L11 ANSWER 14 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1978:51165 CAPLUS

DOCUMENT NUMBER: 88:51165

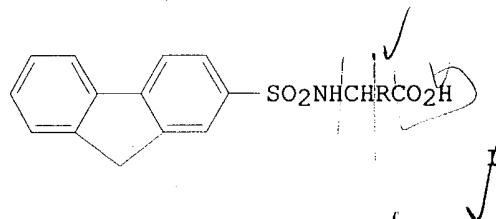
TITLE: Pharmaceutical N-sulfonylaminocarboxylic acids

INVENTOR(S): Toyoshima, Shigehsi; Kanao, Seizo; Toyoda, Takeshi;
 Suyama, Tadashi

PATENT ASSIGNEE(S): Ajinomoto Co., Inc., Japan
 SOURCE: Ger. Offen., 10 pp. Division of Ger. Offen. 2,043,933.
 CODEN: GWXXBX
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 4
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 2065966	B2	19780302	DE 1970-2065966	19700904
DE 2065966	C3	19781116		
NL 7013043	A	19710309	NL 1970-13043	19700903
US 3801633	A	19740402	US 1970-69993	19700904
GB 1269908	A	19720406	GB 1970-1269908	19700907
GB 1288020	A	19720906	GB 1970-1288020	19700907
US 3850968	A	19741126	US 1973-340644	19730313
PRIORITY APPLN. INFO.:			JP 1969-70716	19690906
			US 1970-69993	19700904

ED Entered STN: 12 May 1984
 GI



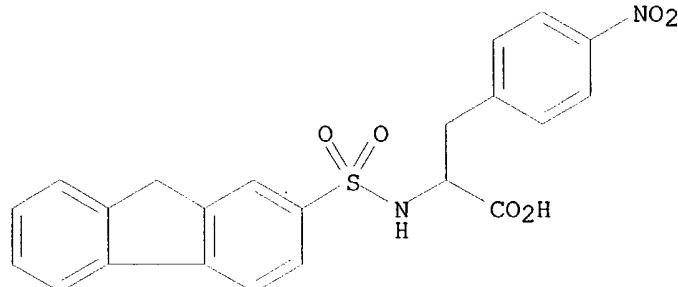
AB Fluorene derivs. I [R = Me (L), CHMe2(L), 4-O2NC6H4CH2 (DL)], useful against influenza virus (extensive data given), were prep'd. by Schotten-Baumann reaktion of L-alanine, L-valine, or DL-4-O2NC6H4CH2CH(NH2)CO2H with 2-fluorenensulfonyl chloride. Also prep'd. were the Na, K, NH4, and HOCH2CH2NH2 salts of L-I (R = Me).

IT 32925-03-4P 32945-11-2P 56211-81-5P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)
 (prep'n. and virucidal activity of)

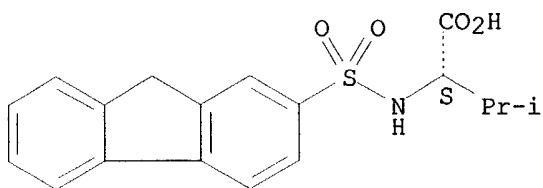
RN 32925-03-4 CAPLUS

CN Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-4-nitro- (9CI) (CA INDEX NAME)



RN 32945-11-2 CAPLUS
 CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

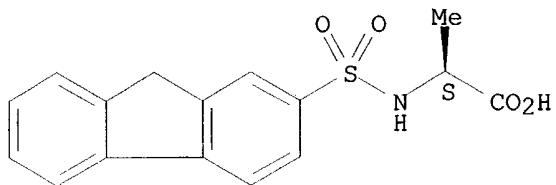
Absolute stereochemistry.



RN 56211-81-5 CAPLUS

CN L-Alanine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



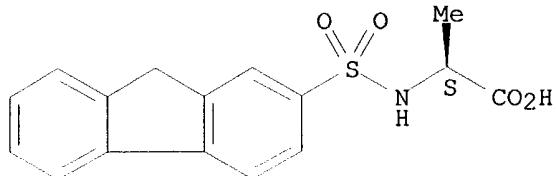
IT 65175-43-1P 65175-44-2P 65175-45-3P

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

RN 65175-43-1 CAPLUS

CN L-Alanine, N-(9H-fluoren-2-ylsulfonyl)-, monosodium salt (9CI) (CA INDEX NAME)

Absolute stereochemistry.

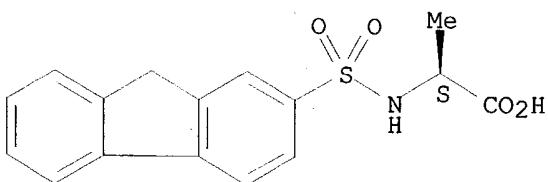


● Na

RN 65175-44-2 CAPLUS

CN L-Alanine, N-(9H-fluoren-2-ylsulfonyl)-, monoammonium salt (9CI) (CA INDEX NAME)

Absolute stereochemistry.



● NH₃

RN 65175-45-3 CAPLUS

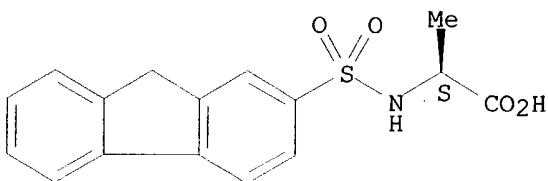
CN L-Alanine, N-(9H-fluoren-2-ylsulfonyl)-, compd. with 2-aminoethanol (1:1)
(9CI) (CA INDEX NAME)

CM 1

CRN 56211-81-5

CMF C16 H15 N O4 S

Absolute stereochemistry.



CM 2

CRN 141-43-5

CMF C2 H7 N O

H₂N—CH₂—CH₂—OH

L11 ANSWER 15 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1977:30073 CAPLUS

DOCUMENT NUMBER:

86:30073

TITLE:

Amino acid derivatives

INVENTOR(S):

Toyoshima, Shigehsi; Kanao, Seizo; Toyoda, Takeshi;

PATENT ASSIGNEE(S):

Suyama, Tadashi; Shimizu, Akira

SOURCE:

Ajinomoto Co., Inc., Japan

Jpn. Kokai Tokkyo Koho, 5 pp. Division of Japan. Kokai

73 28,612.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.

JP 51105048

KIND DATE

A2 19760917

APPLICATION NO. DATE

JP 1976-14276 19760212

Searched by Barb O'Bryen, STIC 571-272-2518

JP 55004740 B4 19800131

PRIORITY APPLN. INFO.:

JP 1976-14276

19760212

ED Entered STN: 12 May 1984

AB Amino acid derivs. were prep'd. by treating phenylalanine or tryptophan with .beta.-naphthalenesulfonyl halides in the presence of alkali or by treating alanine, isoleucine, tryptophan, methionine, or phenylglycine with 2-fluorenesulfonyl halides in the presence of alkali. The products had antiinfluenza viral and anticarcinogenic activities (data given in mice). The LD50 were 750-1500 mg/kg (i.v.) in mice. Thus, 4.5 g .beta.-naphthalenesulfonyl chloride in Et₂O was added to a mixt. of 3.3 g phenylalanine, 10 ml 10% aq. NaOH, and 50 ml 10% aq. Na₂CO₃ over 20 min at room temp. and the mixt. was stirred for 3 h to give 54% N-.beta.-naphthalenesulfonyl-L-phenylalanine. N-.beta.-naphthalenesulfonyl-DL-tryptophan, N-2-fluorenesulfonyl-L-alanine, N-2-fluorenesulfonyl-L-isoleucine, N-2-fluorenesulfonyl-DL-tryptophan, N-2-fluorenesulfonyl-DL-methionine, and N-2-fluorenesulfonyl-DL-phenylglycine were also prep'd.

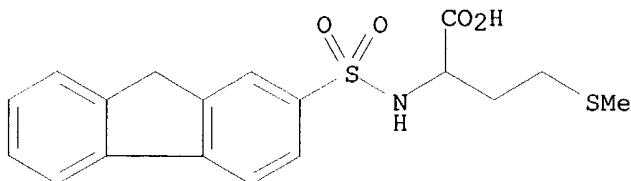
IT 40356-16-9P 56211-81-5P 56211-82-6P

56211-83-7P 61447-77-6P

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. and antiviral and anticarcinogenic activity of)

RN 40356-16-9 CAPLUS

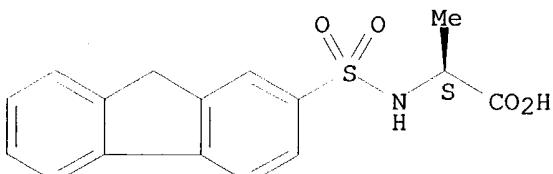
CN Methionine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)



RN 56211-81-5 CAPLUS

CN L-Alanine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

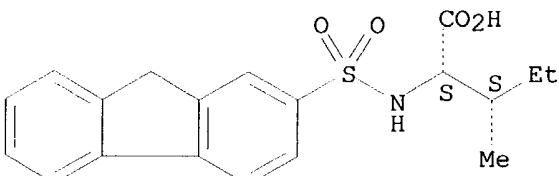
Absolute stereochemistry.



RN 56211-82-6 CAPLUS

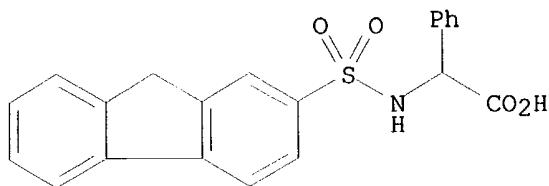
CN L-Isoleucine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



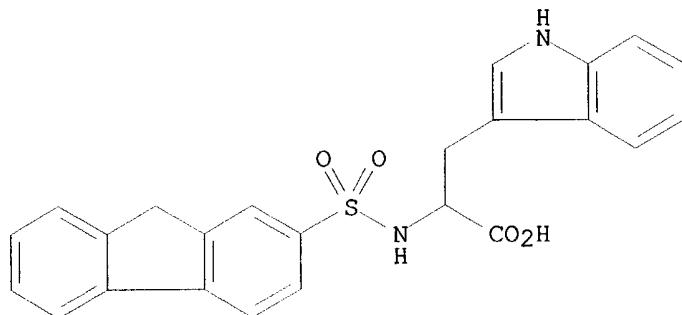
RN 56211-83-7 CAPLUS

CN Benzeneacetic acid, .alpha.-[(9H-fluoren-2-ylsulfonyl)amino]- (9CI) (CA INDEX NAME)



RN 61447-77-6 CAPLUS

CN Tryptophan, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)



L11 ANSWER 16 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1974:83639 CAPLUS

DOCUMENT NUMBER:

80:83639

TITLE:

N-2-Fluorenesulfonylamino acids

INVENTOR(S):

Toshima, Shigeru; Toyota, Takeshi; Suyama, Tadashi

PATENT ASSIGNEE(S):

Ajinomoto Co., Inc.

SOURCE:

Jpn. Tokkyo Koho, 3 pp.

CODEN: JAXXAD

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 48039932	B4	19731128	JP 1970-100902	19701116
PRIORITY APPLN. INFO.:			JP 1970-100902	19701116

ED Entered STN: 12 May 1984

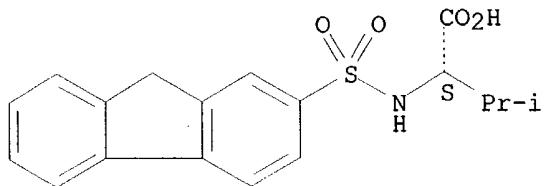
AB Condensation of 2-fluorenesulfonyl chloride (I) with .beta.-alanine (II), L-valine (III), or 4-nitrophenylalanine (IV) in the presence of an alkali gave the corresponding title compds., which have anti-influenza activity and low toxicity. Thus, a mixt. of 20 g 2-fluorenesulfonic acid, 35 g PC15, and 90 ml PC13 was refluxed 1 hr in a water bath and the reaction mixt. poured into ice-water to give 16.5 g I. To a soln. of 1.8 g II in 10% NaOH were added 5.3 g I in acetone and 5.3 g of Na2CO3 in H2O, alternately, at room temp. with stirring in 30 min and the soln. stirred 30 min at 50.degree. and 3 hr at room temp. to give 87% N-2-fluorenesulfonylalanine. Similarly, III and IV gave 70 and 57%, resp., of the corresponding condensates.

IT 32945-11-2P 52525-95-8P

RL: SPN (Synthetic preparation); PREP (Preparation)
(prep. of)

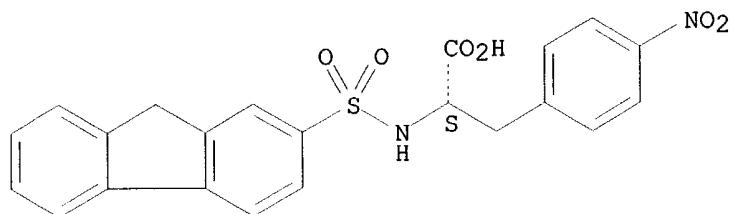
RN 32945-11-2 CAPLUS
 CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 52525-95-8 CAPLUS
 CN L-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-4-nitro- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

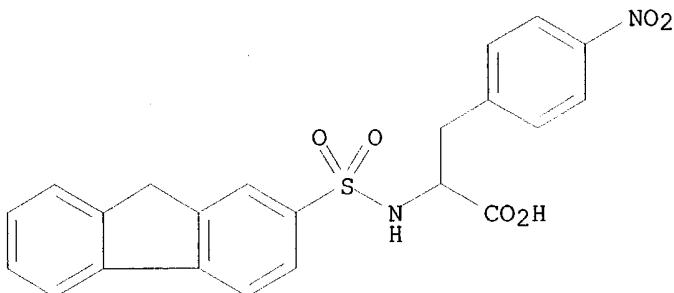


L11 ANSWER 17 OF 25 CAPLUS COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 1971:421010 CAPLUS
 DOCUMENT NUMBER: 75:21010
 TITLE: Antiviral N-acyl-, sulfonyl-, and alkylamino acids
 INVENTOR(S): Toyoshima, Shigeshi; Kanao, Saizu; Toyoda, Takeshi;
 Suyama, Tadashi
 PATENT ASSIGNEE(S): Ajinomoto Co. Inc.
 SOURCE: Ger. Offen., 21 pp.
 CODEN: GWXXBX
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 4
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 2043933	A	19710311	DE 1970-2043933	19700904
DE 2043933	B2	19770908		
NL 7013043	A	19710309	NL 1970-13043	19700903
US 3801633	A	19740402	US 1970-69993	19700904
GB 1269908	A	19720406	GB 1970-1269908	19700907
GB 1288020	A	19720906	GB 1970-1288020	19700907
US 3850968	A	19741126	US 1973-340644	19730313
PRIORITY APPLN. INFO.:			JP 1969-70716	19690906
			US 1970-69993	19700904

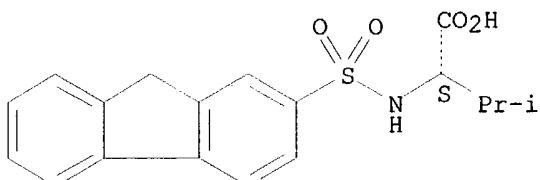
ED Entered STN: 12 May 1984
 AB The title compds., effective against influenza A-2 and type B Lee strain viruses, were prep'd. Thus, α -naphthylacetic acid and PC13 gave α -naphthylacetyl chloride (I). Reaction of L-leucine with I in aq. NaOH-NaHCO₃ yielded 78% N-(α -naphthylacetyl)-L-leucine. Among 13 title compds. similarly prep'd. were N-(carboxymethyl)-L-phenylalanine, N-(p-nitrophenoxyacetyl)-p-nitrophenylalanine, and N-(2-fluorenylsulfonyl)-

.beta.-alanine.
 IT 32925-03-4P 32945-11-2P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (prepn. of).
 RN 32925-03-4 CAPLUS
 CN Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-4-nitro- (9CI) (CA INDEX NAME)



RN 32945-11-2 CAPLUS
 CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L11 ANSWER 18 OF 25 USPATFULL on STN
 ACCESSION NUMBER: 2004:39403 USPATFULL
 TITLE: Method of inhibiting matrix metalloproteinases
 INVENTOR(S): O'Brien, Patrick Michael, Stockbridge, MI, UNITED STATES
 Picard, Joseph Armand, Canton, MI, UNITED STATES
 Sliskovic, Drago Robert, Saline, MI, UNITED STATES
 White, Andrew David, Pinckney, MI, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2004029945	A1	20040212
APPLICATION INFO.:	US 2003-603677	A1	20030625 (10)
RELATED APPLN. INFO.:	Division of Ser. No. US 2002-162518, filed on 4 Jun 2002, GRANTED, Pat. No. US 6620835 Division of Ser. No. US 1999-254384, filed on 2 Mar 1999, GRANTED, Pat. No. US 6624177 A 371 of International Ser. No. WO 1997-US14859, filed on 22 Aug 1997, PENDING		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1996-25062P	19960904 (60)
	US 1997-55713P	19970807 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	WARNER-LAMBERT COMPANY, 2800 PLYMOUTH RD, ANN ARBOR, MI, 48105	

NUMBER OF CLAIMS: 3

EXEMPLARY CLAIM: 1

LINE COUNT: 1256

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to a method of inhibiting matrix metalloproteinases using compounds that are dibenzofuran sulfonamide derivatives having the Formula I ##STR1##

More particularly, the present invention relates to a method of treating diseases in which matrix metalloproteinases are involved such as multiple sclerosis, atherosclerotic plaque rupture, restenosis, aortic aneurism, heart failure, periodontal disease, corneal ulceration, burns, decubital ulcers, chronic ulcers or wounds, cancer metastasis, tumor angiogenesis, arthritis, or other autoimmune or inflammatory diseases dependent upon tissue invasion by leukocytes.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

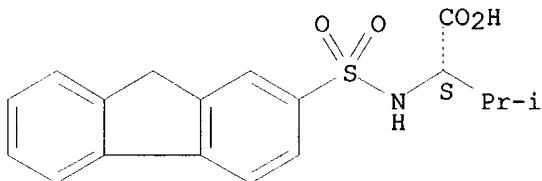
IT 32945-11-2P 204440-69-7P 204440-70-0P
204440-71-1P 204440-91-5P 204440-92-6P
204440-93-7P

(prepn. of dibenzofuransulfonamides as matrix metalloproteinase inhibitors and their therapeutic uses)

RN 32945-11-2 USPATFULL

CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

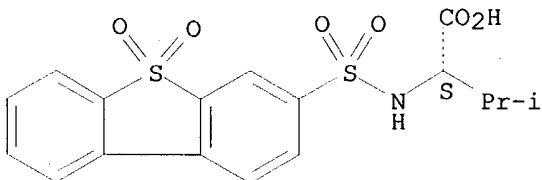
Absolute stereochemistry.



RN 204440-69-7 USPATFULL

CN L-Valine, N-[(5,5-dioxido-3-dibenzothienyl)sulfonyl]- (9CI) (CA INDEX NAME)

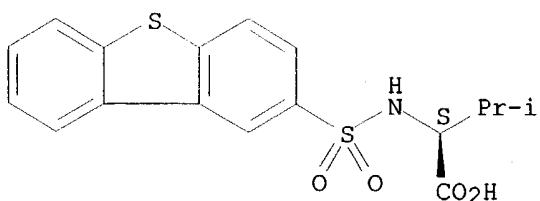
Absolute stereochemistry.



RN 204440-70-0 USPATFULL

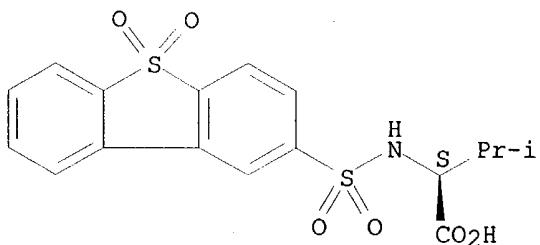
CN L-Valine, N-(2-dibenzothienylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



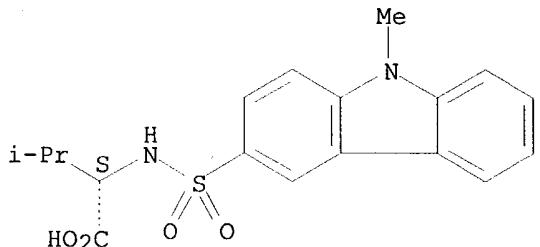
RN 204440-71-1 USPATFULL
 CN L-Valine, N-[(5,5-dioxido-2-dibenzothienyl)sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



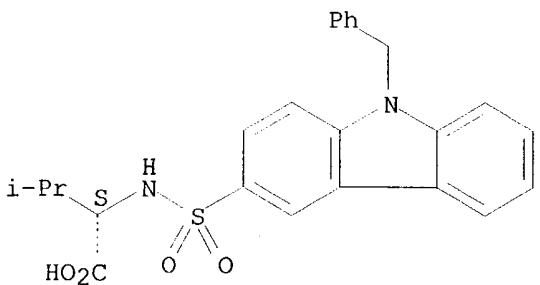
RN 204440-91-5 USPATFULL
 CN L-Valine, N-[(9-methyl-9H-carbazol-3-yl)sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



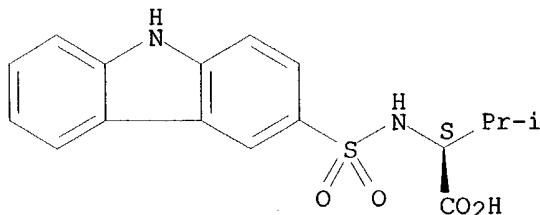
RN 204440-92-6 USPATFULL
 CN L-Valine, N-[(9-(phenylmethyl)-9H-carbazol-3-yl)sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 204440-93-7 USPATFULL
 CN L-Valine, N-(9H-carbazol-3-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L11 ANSWER 19 OF 25 USPATFULL on STN

ACCESSION NUMBER: 2003:45353 USPATFULL

TITLE: Method of inhibiting matrix metalloproteinases

INVENTOR(S): O'Brien, Patrick Michael, Stockbridge, MI, UNITED STATES

Picard, Joseph Armand, Canton, MI, UNITED STATES

Sliskovic, Drago Robert, Saline, MI, UNITED STATES

White, Andrew David, Pinckney, MI, UNITED STATES

PATENT INFORMATION:

	NUMBER	KIND	DATE
US 2003092665		A1	20030213
US 6620835		B2	20030916
US 2002-162518		A1	20020604 (10)

APPLICATION INFO.:

RELATED APPLN. INFO.: Division of Ser. No. US 1999-254384, filed on 2 Mar 1999, PENDING A 371 of International Ser. No. WO 1997-US14859, filed on 22 Aug 1997, UNKNOWN

PRIORITY INFORMATION:

	NUMBER	DATE
US 1996-25062P		19960904 (60)
US 1997-55713P		19970807 (60)

DOCUMENT TYPE:

Utility

FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE:

Andrew J. Leon, Warner-Lambert Company, 2800 Plymouth Road, Ann Arbor, MI, 48105

NUMBER OF CLAIMS:

33

EXEMPLARY CLAIM:

1

LINE COUNT:

1445

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to a method of inhibiting matrix metalloproteinases using compounds that are dibenzofuran sulfonamide derivatives having the Formula I ##STR1##

More particularly, the present invention relates to a method of treating diseases in which matrix metalloproteinases are involved such as multiple sclerosis, atherosclerotic plaque rupture, restenosis, aortic aneurism, heart failure, periodontal disease, corneal ulceration, burns, decubital ulcers, chronic ulcers or wounds, cancer metastasis, tumor angiogenesis, arthritis, or other autoimmune or inflammatory diseases dependent upon tissue invasion by leukocytes.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

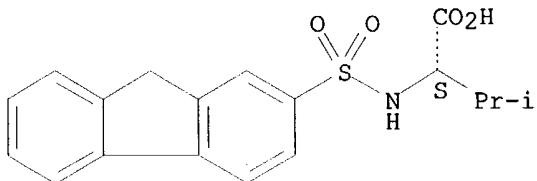
IT 32945-11-2P 204440-69-7P 204440-70-0P
204440-71-1P 204440-91-5P 204440-92-6P
204440-93-7P

(prep. of dibenzofuransulfonamides as matrix metalloproteinase inhibitors and their therapeutic uses)

RN 32945-11-2 USPATFULL

CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

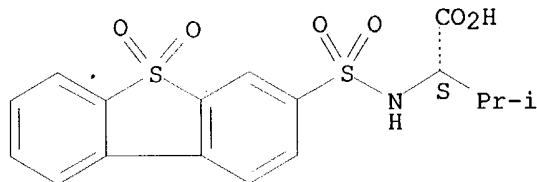
Absolute stereochemistry.



RN 204440-69-7 USPATFULL

CN L-Valine, N-[(5,5-dioxido-3-dibenzothienyl)sulfonyl]- (9CI) (CA INDEX NAME)

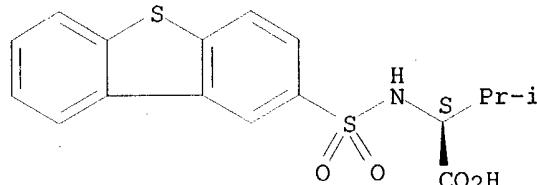
Absolute stereochemistry.



RN 204440-70-0 USPATFULL

CN L-Valine, N-(2-dibenzothienylsulfonyl)- (9CI) (CA INDEX NAME)

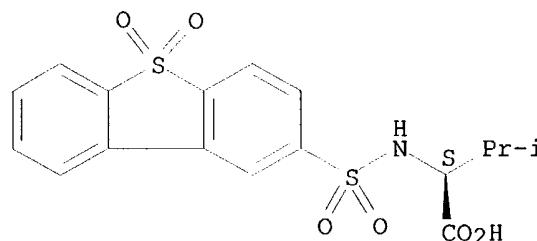
Absolute stereochemistry.



RN 204440-71-1 USPATFULL

CN L-Valine, N-[(5,5-dioxido-2-dibenzothienyl)sulfonyl]- (9CI) (CA INDEX NAME)

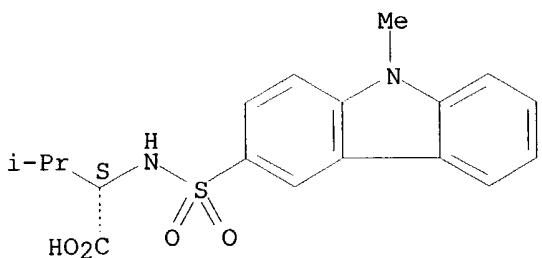
Absolute stereochemistry.



RN 204440-91-5 USPATFULL

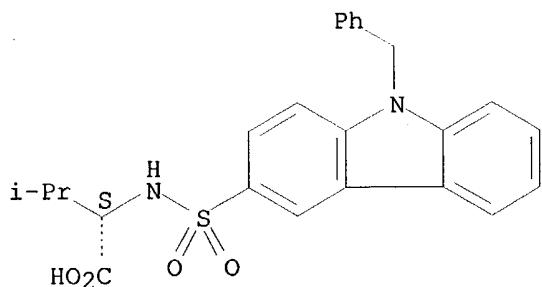
CN L-Valine, N-[(9-methyl-9H-carbazol-3-yl)sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



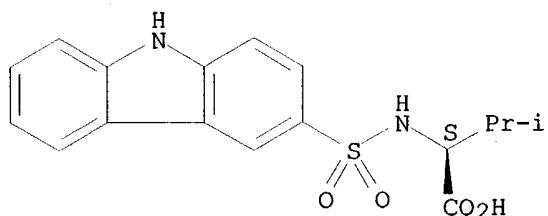
RN 204440-92-6 USPATFULL
 CN L-Valine, N-[9-(phenylmethyl)-9H-carbazol-3-yl]sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 204440-93-7 USPATFULL
 CN L-Valine, N-(9H-carbazol-3-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L11 ANSWER 20 OF 25 USPATFULL on STN
 ACCESSION NUMBER: 2003:253637 USPATFULL
 TITLE: Matrix metalloproteinase inhibitors and their therapeutic uses
 INVENTOR(S): O'Brien, Patrick Michael, Stockbridge, MI, United States
 Picard, Joseph Armand, Canton, MI, United States
 Sliskovic, Drago Robert, Saline, MI, United States
 White, Andrew David, Pinckney, MI, United States
 PATENT ASSIGNEE(S): Warner-Lambert Company, Morris Plains, NJ, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6624177	B1	20030923
	WO 9809934		19980312
APPLICATION INFO.:	US 1999-254384		19990302 (9)

WO 1997-US14859

19970822

19990302 PCT 371 date

NUMBER DATE

PRIORITY INFORMATION: US 1996-25062P 19960904 (60)
 DOCUMENT TYPE: Utility
 FILE SEGMENT: GRANTED
 PRIMARY EXAMINER: Shah, Mukund J.
 ASSISTANT EXAMINER: McKenzie, Thomas
 LEGAL REPRESENTATIVE: Leon, Andrew J., Crissey, Todd M.
 NUMBER OF CLAIMS: 15
 EXEMPLARY CLAIM: 1
 NUMBER OF DRAWINGS: 0 Drawing Figure(s); 0 Drawing Page(s)
 LINE COUNT: 1249

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to a method of inhibiting matrix metalloproteinases using compounds that are dibenzofuran sulfonamide derivatives having the Formula I ##STR1##

More particularly, the present invention relates to a method of treating diseases in which matrix metalloproteinases are involved such as multiple sclerosis, atherosclerotic plaque rupture, restenosis, aortic aneurism, heart failure, periodontal disease, corneal ulceration, burns, decubital ulcers, chronic ulcers or wounds, cancer metastasis, tumor angiogenesis, arthritis, or other autoimmune or inflammatory diseases dependent upon tissue invasion by leukocytes.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

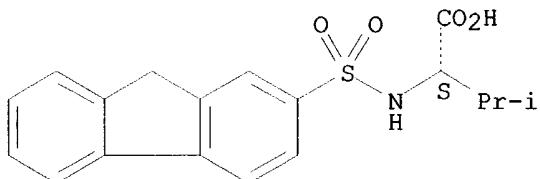
IT 32945-11-2P 204440-69-7P 204440-70-0P
 204440-71-1P 204440-91-5P 204440-92-6P
 204440-93-7P

(prepn. of dibenzofuransulfonamides as matrix metalloproteinase inhibitors and their therapeutic uses)

RN 32945-11-2 USPATFULL

CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

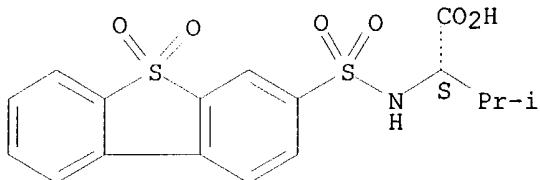
Absolute stereochemistry.



RN 204440-69-7 USPATFULL

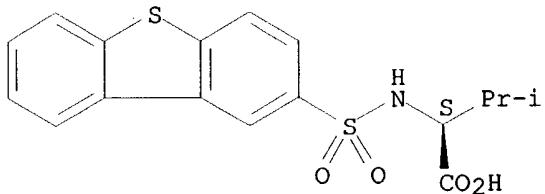
CN L-Valine, N-[(5,5-dioxido-3-dibenzothienyl)sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



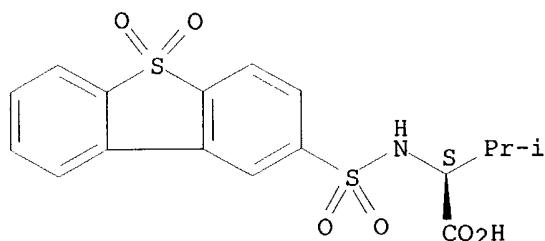
RN 204440-70-0 USPATFULL
 CN L-Valine, N-(2-dibenzothienylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



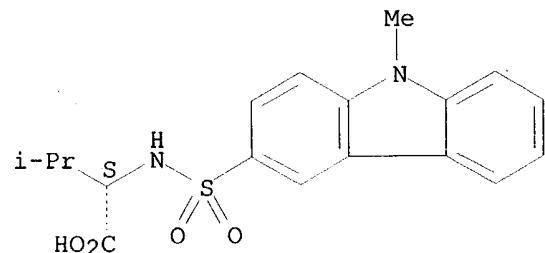
RN 204440-71-1 USPATFULL
 CN L-Valine, N-[(5,5-dioxido-2-dibenzothienyl)sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



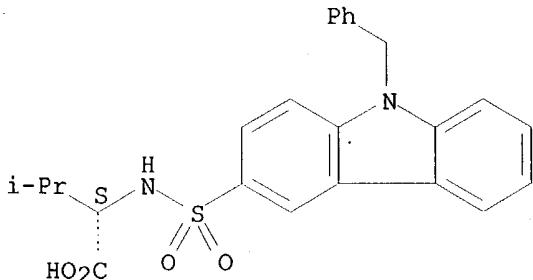
RN 204440-91-5 USPATFULL
 CN L-Valine, N-[(9-methyl-9H-carbazol-3-yl)sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



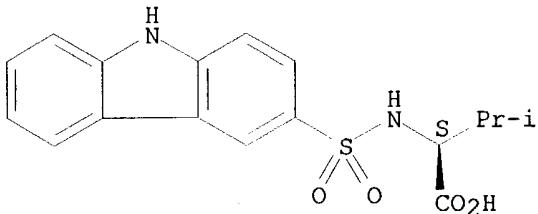
RN 204440-92-6 USPATFULL
 CN L-Valine, N-[[9-(phenylmethyl)-9H-carbazol-3-yl]sulfonyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 204440-93-7 USPATFULL
 CN L-Valine, N-(9H-carbazol-3-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L11 ANSWER 21 OF 25 USPATFULL on STN
 ACCESSION NUMBER: 2001:163339 USPATFULL
 TITLE: Dibenzo-furan sulfonamide matrix metalloproteinase inhibitors
 INVENTOR(S): Picard, Joseph Armand, Canton, MI, United States
 Sliskovic, Drago Robert, Saline, MI, United States
 PATENT ASSIGNEE(S): Warner-Lambert Company, Morris Plains, NJ, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6294674	B1	20010925
	WO 9809957		19980312
APPLICATION INFO.:	US 1999-254403		19990302 (9)
	WO 1997-US15444		19970902
			19990302 PCT 371 date
			19990302 PCT 102(e) date

	NUMBER	DATE
PRIORITY INFORMATION:	US 1996-25063P	19960904 (60)
	US 1997-55714P	19970807 (60)

DOCUMENT TYPE: Utility
 FILE SEGMENT: GRANTED
 PRIMARY EXAMINER: Stockton, Laura L.
 LEGAL REPRESENTATIVE: Ashbrook, Charles W.
 NUMBER OF CLAIMS: 1
 EXEMPLARY CLAIM: 1
 LINE COUNT: 1871
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to compounds of Formula I that inhibit matrix metalloproteinases and to a method of inhibiting matrix metalloproteinases using the compounds. ##STR1##

wherein Q is an un-natural amino acid. More particularly, the present invention relates to a method of treating diseases in which matrix metalloproteinases are involved such as multiple sclerosis, atherosclerotic plaque rupture, restenosis, aortic aneurism, heart failure, periodontal disease, corneal ulceration, burns, decubital ulcers, chronic ulcers or wounds, cancer metastasis, tumor angiogenesis, arthritis, or other autoimmune or inflammatory diseases dependent upon tissue invasion by leukocytes.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

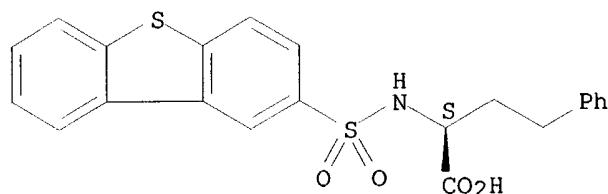
IT 204769-50-6P 204769-91-5P 204769-92-6P

(prepn. of dibenzofuransulfonyl and related amino acids for inhibition of matrix metalloproteinases)

RN 204769-50-6 USPATFULL

CN Benzenebutanoic acid, .alpha.-[(2-dibenzothienylsulfonyl)amino]-, (S)-
(9CI) (CA INDEX NAME)

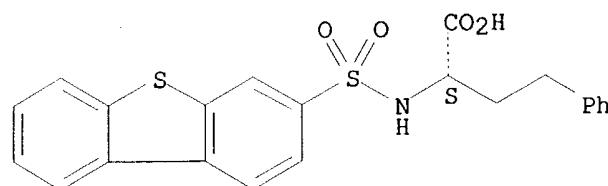
Absolute stereochemistry.



RN 204769-91-5 USPATFULL

CN Benzenebutanoic acid, .alpha.-[(3-dibenzothienylsulfonyl)amino]-, (S)-
(9CI) (CA INDEX NAME)

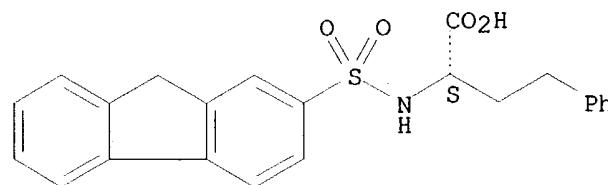
Absolute stereochemistry.



RN 204769-92-6 USPATFULL

CN Benzenebutanoic acid, .alpha.-[(9H-fluoren-2-ylsulfonyl)amino]-,
(.alpha.S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L11 ANSWER 22 OF 25 USPATFULL on STN

ACCESSION NUMBER: 1999:106456 USPATFULL

TITLE: Method for treating and preventing heart failure and ventricular dilatation

INVENTOR(S): Peterson, Jr., Joseph Thomas, Brighton, MI, United States
 PATENT ASSIGNEE(S): Pressler, Milton Lethan, Saline, MI, United States
 Warner-Lambert Company, Morris Plains, NJ, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5948780		19990907
APPLICATION INFO.:	US 1997-987167		19971208 (8)

	NUMBER	DATE
PRIORITY INFORMATION:	US 1996-32631P	19961209 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	Granted	
PRIMARY EXAMINER:	Henley, III, Raymond	
LEGAL REPRESENTATIVE:	Ashbrook, Charles W.	
NUMBER OF CLAIMS:	21	
EXEMPLARY CLAIM:	1	
LINE COUNT:	4703	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Matrix metalloproteinase inhibitors are useful for preventing and treating heart failure, and ventricular dilatation in mammals.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

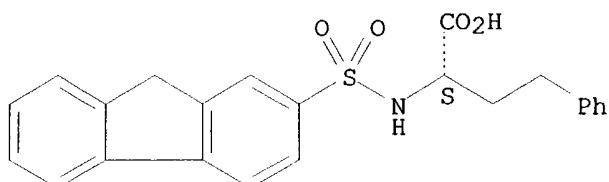
IT 204769-92-6

(use of matrix metalloproteinase inhibitors in treating heart failure and ventricular dilation)

RN 204769-92-6 USPATFULL

CN Benzenebutanoic acid, .alpha.-[(9H-fluoren-2-ylsulfonyl)amino]-, (.alpha.S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L11 ANSWER 23 OF 25 USPATFULL on STN
 ACCESSION NUMBER: 75:61131 USPATFULL
 TITLE: N-ethylcarbaminomethylisoleucine
 INVENTOR(S): Toyoshima, Shigeshi, Tokyo, Japan
 Kanao, Seizo, Tokyo, Japan
 Toyoda, Takeshi, Sagamihara, Japan
 Suyama, Tadashi, Kawasaki, Japan
 PATENT ASSIGNEE(S): Ajinomoto Co., Inc., Tokyo, Japan (non-U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 3919291		19751111
APPLICATION INFO.:	US 1974-451209		19740314 (5)
RELATED APPLN. INFO.:	Division of Ser. No. US 1972-281042, filed on 16 Aug 1972, now patented, Pat. No. US 3845097 which is a continuation-in-part of Ser. No. US 1970-69993, filed on 4 Sep 1970, now patented, Pat. No. US 3801633		

	NUMBER	DATE
PRIORITY INFORMATION:	JP 1971-63250	19710819
	JP 1971-63252	19710819
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	Granted	
PRIMARY EXAMINER:	Raymond, Richard L.	
LEGAL REPRESENTATIVE:	Berman, Hans	
NUMBER OF CLAIMS:	1	
EXEMPLARY CLAIM:	1	
LINE COUNT:	580	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The following N-substituted amino acids have been found to combat infection with influenza virus, to counteract inflammation, or to have anti-tumor effects in mice while being relatively non-toxic in effective amounts:

N-.beta.-naphthylaminoethylleucine,
N-.beta.-naphthylaminomethylurethane,
N-furfurylaminoethylphenylalanine,
N-furfuryl-4-nitrophenylalanine,
N-benzylvaline,
N-2-fluoresulfonylmethionine,
N-2-fluoresulfonylphenylalanine,
N-lauroylleucine,
N-ethylcarbaminomethylisoleucine,
N-.beta.-naphthylaminomethylthreonine,
N-9-fluorenylacetylphenylalanine,
N-myristoylisoleucine,
N-.beta.-naphthalenesulfonyltryptophan, and
N-propionylvaline.

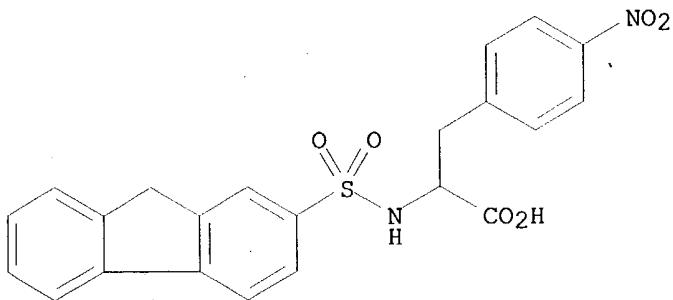
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 32925-03-4P 32945-11-2P

(prepn. of)

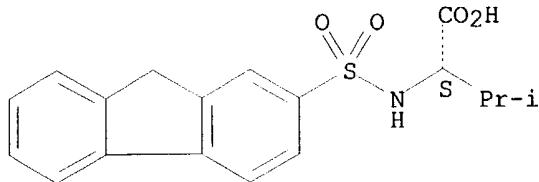
RN 32925-03-4 USPATFULL

CN Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-4-nitro- (9CI) (CA INDEX NAME)



RN 32945-11-2 USPATFULL
 CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L11 ANSWER 24 OF 25 TOXCENTER COPYRIGHT 2004 ACS on STN
 ACCESSION NUMBER: 1977:68779 TOXCENTER
 COPYRIGHT: Copyright 2004 ACS
 DOCUMENT NUMBER: CA08605030073U
 TITLE: Amino acid derivatives
 AUTHOR(S): Toyoshima, Shigeshi; Kanao, Seizo; Toyoda, Takeshi;
 Suyama, Tadashi; Shimizu, Akira
 CORPORATE SOURCE: ASSIGNEE: Ajinomoto Co., Inc.
 PATENT INFORMATION: JP 76105048 17 Sep 1976
 SOURCE: (1976) Jpn. Kokai Tokkyo Koho, 5 pp. Division of Japan.
 Kokai 73 28,612.
 CODEN: JKXXAF.
 COUNTRY: JAPAN
 DOCUMENT TYPE: Patent
 FILE SEGMENT: CAPLUS
 OTHER SOURCE: CAPLUS 1977:30073
 LANGUAGE: Japanese
 ENTRY DATE: Entered STN: 20011116
 Last Updated on STN: 20021210

ABSTRACT:
 Amino acid derivs. were prep'd. by treating phenylalanine or tryptophan with .beta.-naphthalenesulfonyl halides in the presence of alkali or by treating alanine, isoleucine, tryptophan, methionine, or phenylglycine with 2-fluorenesulfonyl halides in the presence of alkali. The products had antiinfluenza viral and anticarcinogenic activities (data given in mice). The LD₅₀ were 750-1500 mg/kg (i.v.) in mice. Thus, 4.5 g .beta.-naphthalenesulfonyl chloride in Et₂O was added to a mixt. of 3.3 g phenylalanine, 10 ml 10% aq. NaOH, and 50 ml 10% aq. Na₂CO₃ over 20 min at room temp. and the mixt. was stirred for 3 h to give 54% N-.beta.-naphthalenesulfonyl-L-phenylalanine. N-.beta.-naphthalenesulfonyl-DL-trypophan, N-2-fluorenesulfonyl-L-alanine, N-2-fluorenesulfonyl-L-isoleucine,

N-2-fluorenesulfonyl-DL-tryptophan, N-2-fluorenesulfonyl-DL-methionine, and N-2-fluorenesulfonyl-DL-phenylglycine were also prep'd.

CLASSIFICATION CODE: 34-2

SUPPLEMENTARY TERMS: Miscellaneous Descriptors

amino acid naphthylsulfonyl fluorenesulfonyl; virucide
naphthylsulfonyl amino acid; neoplasm inhibitor
naphthylsulfonyl amino acid

REGISTRY NUMBER: 63-91-2; 40356-16-9; 40356-23-8; 55953-52-1;
56211-81-5; 56211-82-6;
56211-83-7; 61447-77-6; 93-11-8

L11 ANSWER 25 OF 25 TOXCENTER COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1974:65201 TOXCENTER

COPYRIGHT: Copyright 2004 ACS

DOCUMENT NUMBER: CA08015083639R

TITLE: N-2-Fluorenesulfonylamino acids

AUTHOR(S): Teshima, Shigeru; Toyota, Takeshi; Suyama, Tadashi

CORPORATE SOURCE: ASSIGNEE: Ajinomoto Co., Inc.

PATENT INFORMATION: JP 7339932 28 Nov 1973

SOURCE: (1973) Jpn. Tokkyo Koho, 3 pp.

CODEN: JAXXAD.

DOCUMENT TYPE: Patent

FILE SEGMENT: CAPLUS

OTHER SOURCE: CAPLUS 1974:83639

LANGUAGE: Japanese

ENTRY DATE: Entered STN: 20011116

Last Updated on STN: 20021218

ABSTRACT:

Condensation of 2-fluorenesulfonyl chloride (I) with .beta.-alanine (II), L-valine (III), or 4-nitrophenylalanine (IV) in the presence of an alkali gave the corresponding title compds., which have anti-influenza activity and low toxicity. Thus, a mixt. of 20 g 2-fluorenesulfonic acid, 35 g PC15, and 90 ml PC13 was refluxed 1 hr in a water bath and the reaction mixt. poured into ice-water to give 16.5 g I. To a soln. of 1.8 g II in 10% NaOH were added 5.3 g I in acetone and 5.3 g of Na₂CO₃ in H₂O, alternately, at room temp. with stirring in 30 min and the soln. stirred 30 min at 50.degree. and 3 hr at room temp. to give 87% N-2-fluorenesulfonylalanine. Similarly, III and IV gave 70 and 57%, resp., of the corresponding condensates.

CLASSIFICATION CODE: 34-2

SUPPLEMENTARY TERMS: Miscellaneous Descriptors

amino acid fluorenylsulfonyl influenza

REGISTRY NUMBER: 52525-94-7; 13354-17-1; 32869-90-2; 32945-11-2;

52525-95-8; 107-95-9; 949-99-5; 72-18-4

=> fil reg

FILE 'REGISTRY' ENTERED AT 14:34:09 ON 24 MAR 2004

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.

PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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STRUCTURE FILE UPDATES: 23 MAR 2004 HIGHEST RN 666817-09-0

DICTIONARY FILE UPDATES: 23 MAR 2004 HIGHEST RN 666817-09-0

TSCA INFORMATION NOW CURRENT THROUGH JANUARY 6, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:
<http://www.cas.org/ONLINE/DBSS/registryss.html>

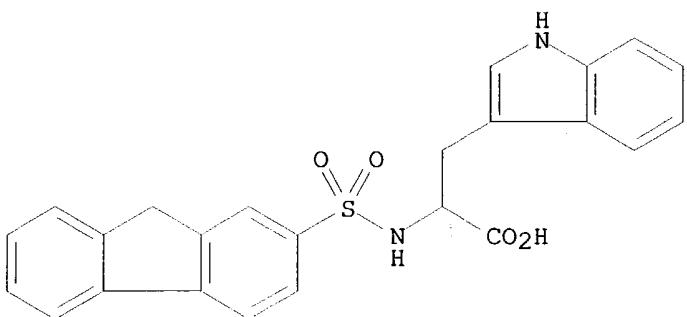
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 61447-77-6 or 40356-16-9

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 (52525-95-8/RN)
 1 56211-81-5
 (56211-81-5/RN)
 1 56211-82-6
 (56211-82-6/RN)
 1 56211-83-7
 (56211-83-7/RN)
 1 61447-77-6
 (61447-77-6/RN)
 1 40356-16-9
 (40356-16-9/RN)

L12 7 32945-11-2 OR 52525-95-8 OR 56211-81-5 OR 56211-82-6 OR
 56211-83-7 OR 61447-77-6 OR 40356-16-9

=> d ide l12 1-7; fil hom

L12 ANSWER 1 OF 7 REGISTRY COPYRIGHT 2004 ACS on STN
 RN 61447-77-6 REGISTRY
 CN Tryptophan, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)
 OTHER CA INDEX NAMES:
 CN DL-Tryptophan, N-(9H-fluoren-2-ylsulfonyl)-
 OTHER NAMES:
 CN N-2-Fluorenesulfonyl-DL-tryptophan
 MF C24 H20 N2 O4 S
 LC STN Files: CA, CAPLUS, TOXCENTER

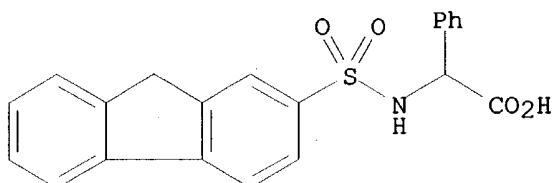


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)
 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L12 ANSWER 2 OF 7 REGISTRY COPYRIGHT 2004 ACS on STN
 RN 56211-83-7 REGISTRY
 CN Benzeneacetic acid, .alpha.-[(9H-fluoren-2-ylsulfonyl)amino]- (9CI) (CA INDEX NAME)
 OTHER CA INDEX NAMES:

CN Benzeneacetic acid, .alpha.-[(9H-fluoren-2-ylsulfonyl)amino]-, (.+-.)-
 OTHER NAMES:
 CN N-2-Fluorenesulfonyl-DL-phenylglycine
 MF C21 H17 N O4 S
 LC STN Files: CA, CAPLUS, TOXCENTER

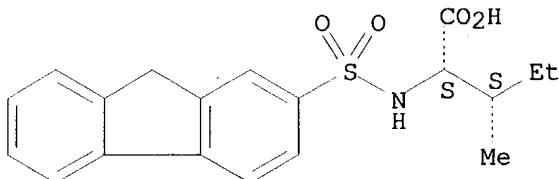


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1907 TO DATE)
 3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L12 ANSWER 3 OF 7 REGISTRY COPYRIGHT 2004 ACS on STN
 RN 56211-82-6 REGISTRY
 CN L-Isoleucine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)
 OTHER NAMES:
 CN N-2-Fluorenesulfonyl-L-isoleucine
 FS STEREOSEARCH
 MF C19 H21 N O4 S
 LC STN Files: CA, CAPLUS, TOXCENTER

Absolute stereochemistry.

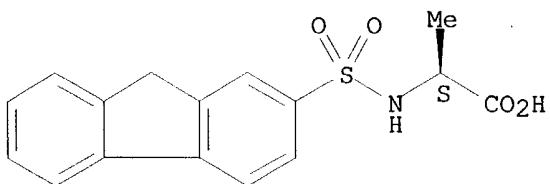


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1907 TO DATE)
 3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L12 ANSWER 4 OF 7 REGISTRY COPYRIGHT 2004 ACS on STN
 RN 56211-81-5 REGISTRY
 CN L-Alanine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)
 OTHER NAMES:
 CN N-2-Fluorenesulfonyl-L-alanine
 FS STEREOSEARCH
 MF C16 H15 N O4 S
 CI COM
 LC STN Files: CA, CAPLUS, CASREACT, TOXCENTER

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

5 REFERENCES IN FILE CA (1907 TO DATE)

5 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L12 ANSWER 5 OF 7 REGISTRY COPYRIGHT 2004 ACS on STN

RN 52525-95-8 REGISTRY

CN L-Phenylalanine, N-(9H-fluoren-2-ylsulfonyl)-4-nitro- (9CI) (CA INDEX NAME)

OTHER NAMES:

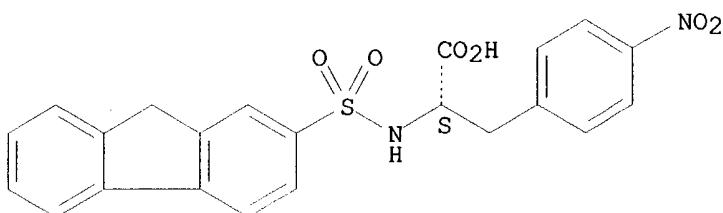
CN N-(Fluoren-2-ylsulfonyl)-3-(4-nitrophenyl)-L-alanine

FS STEREOSEARCH

MF C22 H18 N2 O6 S

LC STN Files: BEILSTEIN*, CA, CAPLUS, CASREACT, TOXCENTER
(*File contains numerically searchable property data)

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)

2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L12 ANSWER 6 OF 7 REGISTRY COPYRIGHT 2004 ACS on STN

RN 40356-16-9 REGISTRY

CN Methionine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

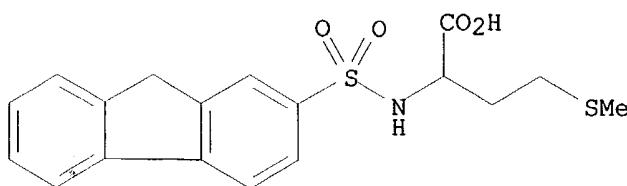
CN DL-Methionine, N-(9H-fluoren-2-ylsulfonyl)-

OTHER NAMES:

CN N-2-Fluorenesulfonyl-DL-methionine

MF C18 H19 N O4 S2

LC STN Files: CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, TOXCENTER

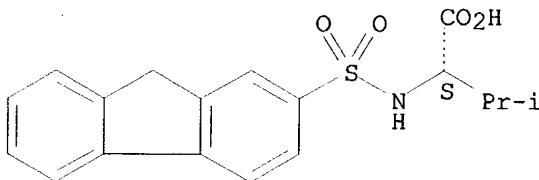


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3 REFERENCES IN FILE CA (1907 TO DATE)
 3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L12 ANSWER 7 OF 7 REGISTRY COPYRIGHT 2004 ACS on STN
 RN 32945-11-2 REGISTRY
 CN L-Valine, N-(9H-fluoren-2-ylsulfonyl)- (9CI) (CA INDEX NAME)
 OTHER CA INDEX NAMES:
 CN Valine, N-(fluoren-2-ylsulfonyl)-, L- (8CI)
 OTHER NAMES:
 CN N-(Fluoren-2-ylsulfonyl)-L-valine
 CN N-2-Fluorenesulfonyl-L-valine
 FS STEREOSEARCH
 MF C18 H19 N O4 S
 LC STN Files: BEILSTEIN*, CA, CAPLUS, CASREACT, TOXCENTER, USPAT2,
 USPATFULL
 (*File contains numerically searchable property data)

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

6 REFERENCES IN FILE CA (1907 TO DATE)
 6 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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L13 0 L7

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